

**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 **September 30, 2012**

OR

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

For the transition period from _____ to _____.

COMMISSION FILE NUMBER: 000-19271

IDEXX LABORATORIES, INC.

(Exact name of registrant as specified in its charter)

DELAWARE

*(State or other jurisdiction of incorporation
or organization)*

01-0393723

(IRS Employer Identification No.)

ONE IDEXX DRIVE, WESTBROOK, MAINE

(Address of principal executive offices)

04092

(ZIP Code)

207-556-0300

(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

Indicate the number of shares outstanding of each of the issuer's classes of common stock, as of the latest practicable date. The number of shares outstanding of the registrant's Common Stock, \$0.10 par value, was 54,672,689 on October 9, 2012.

IDEXX LABORATORIES, INC.
Quarterly Report on Form 10-Q
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PART I— FINANCIAL INFORMATION

Item 1. Financial Statements.

IDEXX LABORATORIES, INC. AND SUBSIDIARIES

CONDENSED CONSOLIDATED BALANCE SHEETS

*(in thousands, except per share amounts)
(Unaudited)*

	<u>September 30,</u> <u>2012</u>	<u>December 31,</u> <u>2011</u>
ASSETS		
Current Assets:		
Cash and cash equivalents	\$ 221,441	\$ 183,895
Accounts receivable, net of reserves of \$3,618 in 2012 and \$3,239 in 2011	141,374	141,275
Inventories	147,474	133,099
Deferred income tax assets	25,430	25,637
Other current assets	<u>35,055</u>	<u>40,321</u>
Total current assets	570,774	524,227
Long-Term Assets:		
Property and equipment, net	231,639	216,777
Goodwill	174,614	172,610
Intangible assets, net	63,363	69,209
Other long-term assets, net	<u>52,961</u>	<u>47,991</u>
Total long-term assets	<u>522,577</u>	<u>506,587</u>
TOTAL ASSETS	<u>\$ 1,093,351</u>	<u>\$ 1,030,814</u>
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current Liabilities:		
Accounts payable	\$ 31,728	\$ 36,551
Accrued liabilities	134,664	141,383
Line of credit	232,000	243,000
Current portion of long-term debt	960	917
Current portion of deferred revenue	<u>16,494</u>	<u>15,028</u>
Total current liabilities	415,846	436,879
Long-Term Liabilities:		
Deferred income tax liabilities	20,650	23,288
Long-term debt, net of current portion	1,776	2,501
Long-term deferred revenue, net of current portion	14,656	10,823
Other long-term liabilities	<u>22,888</u>	<u>17,730</u>
Total long-term liabilities	<u>59,970</u>	<u>54,342</u>
Total liabilities	475,816	491,221
Commitments and Contingencies (Note 13)		
Stockholders' Equity:		
Common stock, \$0.10 par value: Authorized: 120,000 shares; Issued: 99,914 and 99,229 shares in 2012 and 2011, respectively	9,991	9,923
Additional paid-in capital	742,022	702,575
Deferred stock units: Outstanding: 119 units in 2012 and 2011	4,616	4,688
Retained earnings	1,262,239	1,127,326
Accumulated other comprehensive income	14,579	15,443
Treasury stock, at cost: 45,215 and 44,128 shares in 2012 and 2011, respectively	<u>(1,415,940)</u>	<u>(1,320,376)</u>
Total IDEXX Laboratories, Inc. stockholders' equity	617,507	539,579
Noncontrolling interest	<u>28</u>	<u>14</u>
Total stockholders' equity	<u>617,535</u>	<u>539,593</u>
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY	<u>\$ 1,093,351</u>	<u>\$ 1,030,814</u>

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

IDEXX LABORATORIES, INC. AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS
(in thousands, except per share amounts)
(Unaudited)

	For the Three Months Ended		For the Nine Months Ended	
	September 30,		September 30,	
	2012	2011	2012	2011
Revenue:				
Product revenue	\$ 196,900	\$ 191,334	\$ 613,288	\$ 583,221
Service revenue	118,575	109,620	360,512	328,267
Total revenue	315,475	300,954	973,800	911,488
Cost of Revenue:				
Cost of product revenue	72,738	76,831	226,420	227,993
Cost of service revenue	72,102	65,456	217,282	195,870
Total cost of revenue	144,840	142,287	443,702	423,863
Gross profit	170,635	158,667	530,098	487,625
Expenses:				
Sales and marketing	52,067	50,682	164,238	152,641
General and administrative	35,307	32,483	105,760	98,219
Research and development	20,349	19,406	60,964	55,839
Income from operations	62,912	56,096	199,136	180,926
Interest expense	(873)	(928)	(2,992)	(2,438)
Interest income	473	450	1,389	1,238
Income before provision for income taxes	62,512	55,618	197,533	179,726
Provision for income taxes	19,639	17,122	62,606	55,970
Net income	42,873	38,496	134,927	123,756
Less: Net income (loss) attributable to noncontrolling interest	20	(11)	14	(20)
Net income attributable to IDEXX Laboratories, Inc. stockholders	\$ 42,853	\$ 38,507	\$ 134,913	\$ 123,776
Earnings per Share:				
Basic	\$ 0.78	\$ 0.68	\$ 2.45	\$ 2.17
Diluted	\$ 0.76	\$ 0.66	\$ 2.40	\$ 2.11
Weighted Average Shares Outstanding:				
Basic	54,938	56,699	55,074	57,141
Diluted	56,088	58,007	56,270	58,636

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

IDEXX LABORATORIES, INC. AND SUBSIDIARIES

CONDENSED CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME

(in thousands, except per share amounts)
(Unaudited)

	For the Three Months Ended		For the Nine Months Ended	
	September 30,		September 30,	
	2012	2011	2012	2011
Net income	\$ 42,873	\$ 38,496	\$ 134,927	\$ 123,756
Other comprehensive income, net of tax:				
Foreign currency translation adjustments	9,221	(12,780)	4,548	799
Unrealized gain (loss) on investments, net of tax expense (benefit) of \$33 and \$66 in 2012 and (\$145) and (\$119) in 2011	56	(246)	111	(201)
Unrealized (loss) gain on derivative instruments:				
Unrealized (loss) gain, net of tax (benefit) expense of (\$1,294) and (\$1,235) in 2012 and \$1,781 and \$188 in 2011	(2,684)	4,075	(2,329)	541
Less: reclassification adjustment for (gains) losses included in net income, net of tax (expense) benefit of (\$395) and (\$1,434) in 2012 and \$565 and \$1,930 in 2011	(872)	1,192	(3,194)	4,103
Unrealized (loss) gain on derivative instruments	(3,556)	5,267	(5,523)	4,644
Other comprehensive income (loss), net of tax	5,721	(7,759)	(864)	5,242
Comprehensive income	48,594	30,737	134,063	128,998
Less: comprehensive income (loss) attributable to noncontrolling interest	20	(11)	14	(20)
Comprehensive income attributable to IDEXX Laboratories, Inc.	<u>\$ 48,574</u>	<u>\$ 30,748</u>	<u>\$ 134,049</u>	<u>\$ 129,018</u>

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

IDEXX LABORATORIES, INC. AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS
(in thousands)
(Unaudited)

	For the Nine Months Ended	
	September 30,	
	2012	2011
Cash Flows from Operating Activities:		
Net income	\$ 134,927	\$ 123,756
Adjustments to reconcile net income to net cash provided by operating activities:		
Depreciation and amortization	38,490	35,388
Loss on disposal of property and equipment	194	481
Increase (decrease) in deferred compensation liability	176	(320)
Provision for uncollectible accounts	837	997
(Benefit of) provision for deferred income taxes	(1,861)	598
Share-based compensation expense	11,684	11,497
Tax benefit from share-based compensation arrangements	(10,182)	(14,009)
Changes in assets and liabilities:		
Accounts receivable	255	(16,181)
Inventories	(18,078)	(9,732)
Other assets	539	3,056
Accounts payable	(4,893)	8,305
Accrued liabilities	(3,751)	16,664
Deferred revenue	5,151	3,018
Net cash provided by operating activities	<u>153,488</u>	<u>163,518</u>
Cash Flows from Investing Activities:		
Purchases of property and equipment	(43,230)	(39,927)
Proceeds from disposition of pharmaceutical product lines	3,000	3,000
Proceeds from sale of property and equipment	45	223
Acquisitions of intangible asset	(900)	-
Acquisition of a business, net of cash acquired	-	(2,600)
Net cash used by investing activities	<u>(41,085)</u>	<u>(39,304)</u>
Cash Flows from Financing Activities:		
(Payments) borrowings on revolving credit facilities, net	(11,000)	24,903
Payment of notes payable	(682)	(643)
Repurchases of common stock	(91,152)	(166,016)
Proceeds from exercises of stock options and employee stock purchase plans	17,156	26,080
Tax benefit from share-based compensation arrangements	10,182	14,009
Net cash used by financing activities	<u>(75,496)</u>	<u>(101,667)</u>
Net effect of changes in exchange rates on cash	639	2,037
Net increase in cash and cash equivalents	37,546	24,584
Cash and cash equivalents at beginning of period	183,895	156,915
Cash and cash equivalents at end of period	<u>\$ 221,441</u>	<u>\$ 181,499</u>

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

IDEXX LABORATORIES, INC. AND SUBSIDIARIES

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

(Unaudited)

NOTE 1. BASIS OF PRESENTATION AND PRINCIPLES OF CONSOLIDATION

The accompanying condensed consolidated financial statements of IDEXX Laboratories, Inc. (“IDEXX,” the “Company,” “we” or “our”) have been prepared in accordance with accounting principles generally accepted in the United States of America (“U.S. GAAP”) for interim financial information and with the requirements of Regulation S-X, Rule 10-01 for financial statements required to be filed as a part of Form 10-Q.

The accompanying condensed consolidated financial statements include the accounts of IDEXX Laboratories, Inc. and our wholly-owned and majority-owned subsidiaries. All intercompany transactions and balances have been eliminated in consolidation.

The accompanying condensed consolidated financial statements reflect, in the opinion of our management, all adjustments necessary for a fair statement of our financial position and results of operations. All such adjustments are of a recurring nature. The consolidated balance sheet data at December 31, 2011 was derived from audited financial statements, but does not include all disclosures required by U.S. GAAP. The results of operations for the three and nine months ended September 30, 2012 are not necessarily indicative of the results to be expected for the full year or any future period. These condensed consolidated financial statements should be read in conjunction with this Quarterly Report on Form 10-Q for the quarter ended September 30, 2012 and our Annual Report on Form 10-K for the year ended December 31, 2011 filed with the Securities and Exchange Commission.

NOTE 2. ACCOUNTING POLICIES

Significant Accounting Policies

The significant accounting policies used in preparation of these condensed consolidated financial statements for the nine months ended September 30, 2012 are consistent with those discussed in Note 2 to the consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2011, except for our significant accounting policies related to revenue recognition. Effective January 1, 2012, revenue from substantially all U.S. distributors is recognized upon delivery to the distributor because title and risk of loss remains with IDEXX until the product is delivered. Prior to January 1, 2012, we recognized revenue at the time of shipment to U.S. distributors because title and risk of loss passed to the distributors on delivery to the common carrier. This change did not have a material impact on our financial statements.

New Accounting Pronouncements Adopted

In September 2011, the Financial Accounting Standards Board (“FASB”) issued an amendment to the accounting guidance for goodwill in order to simplify how companies test goodwill for impairment. The amendment permits an entity to first assess the qualitative factors to determine whether it is more likely than not that the fair value of the reporting unit is less than its carrying amount as a basis for determining whether it is necessary to perform the two-step goodwill impairment test. The more-likely-than-not threshold is defined as having a likelihood of more than 50 percent. If, after assessing the totality of events or circumstances, an entity determines it is not more likely than not that the fair value of a reporting unit is less than its carrying amount, then performing the two-step impairment test is unnecessary. The amendment is effective for annual and interim goodwill impairment tests performed for fiscal years beginning after December 15, 2011. The adoption of this accounting pronouncement did not have a material impact on our financial statements and we do not expect it to have a material impact on our annual goodwill impairment assessment in the fourth quarter.

In June 2011, the FASB issued an amendment to the accounting guidance for presentation of comprehensive income. Under the amended guidance, an entity may present the total of comprehensive income, the components of net income, and the components of other comprehensive income either in a single continuous statement of comprehensive income or in two separate but consecutive statements. In either case, an entity is required to present each component of net income along with total net income, each component of other comprehensive income along with a total for other comprehensive income, and a total amount for comprehensive income. For public companies, the amendment is effective for fiscal years, and interim periods within those years, beginning after December 15, 2011, and shall be applied retrospectively. Other than a change in presentation, the implementation of this accounting pronouncement did not have a material impact on our financial statements.

In May 2011, the FASB issued an amendment to the accounting guidance for fair value measurement and disclosure. Among other things, the guidance expands the disclosure requirements around fair value measurements categorized in Level 3 of the fair value hierarchy and requires disclosure of the level in the fair value hierarchy of items that are not measured at fair value in the statement of financial position but whose fair value must be disclosed. It also clarifies and expands upon existing requirements for measurement of the fair value of financial assets and liabilities as well as instruments classified in shareholders' equity. The guidance is effective for interim and annual periods beginning after December 15, 2011. The adoption of this accounting pronouncement did not have a material impact on our financial statements.

New Accounting Pronouncements Not Yet Adopted

In December 2011, the FASB issued an amendment to the accounting guidance for disclosure of offsetting assets and liabilities and related arrangements. The amendment expands the disclosure requirements in that entities will be required to disclose both gross information and net information about both instruments and transactions eligible for offset in the statement of financial position and instruments and transactions subject to an agreement similar to a master netting arrangement. The amendment is effective for fiscal years, and interim periods within those years, beginning on or after January 1, 2013, and shall be applied retrospectively. We do not expect the adoption of this accounting pronouncement to have a material impact on our financial statements.

NOTE 3. SHARE-BASED COMPENSATION

The fair value of options, restricted stock units, deferred stock units and employee stock purchase rights awarded during the three and nine months ended September 30, 2012 totaled \$1.1 million and \$17.9 million, respectively, compared to \$0.2 million and \$23.8 million for the three and nine months ended September 30, 2011, respectively.

The total unrecognized compensation expense, net of estimated forfeitures, for unvested share-based compensation awards outstanding at September 30, 2012 was \$36.3 million, which will be recognized over a weighted average period of approximately 1.8 years.

We determine the assumptions used in the valuation of option awards as of the date of grant. Differences in the stock price volatility, expected term or risk-free interest rate may necessitate distinct valuation assumptions at each grant date. As such, we may use different assumptions for options granted throughout the year. Option awards are granted with an exercise price equal to the closing market price of our common stock at the date of grant. We have never paid any cash dividends on our common stock and we have no present intention to pay a dividend; therefore, we assume that no dividends will be paid over the expected terms of option awards. The weighted averages of the valuation assumptions used to determine the fair value of each option award on the date of grant and the weighted average estimated fair values were as follows:

	For the Nine Months Ended	
	September 30,	
	2012	2011
Expected stock price volatility	34 %	33 %
Expected term, in years	4.6	4.8
Risk-free interest rate	0.8 %	2.3 %
Weighted average fair value of options granted	\$ 26.37	\$ 24.87

NOTE 4. INVENTORIES

Inventories include material, labor and overhead, and are stated at the lower of cost (first-in, first-out) or market. The components of inventories were as follows (*in thousands*):

	September 30, 2012	December 31, 2011
Raw materials	\$ 30,873	\$ 28,338
Work-in-process	16,852	14,892
Finished goods	99,749	89,869
	<u>\$ 147,474</u>	<u>\$ 133,099</u>

NOTE 5. GOODWILL AND INTANGIBLE ASSETS, NET

The increase in goodwill during the nine months ended September 30, 2012 resulted from changes in foreign currency exchange rates. The decrease in intangible assets other than goodwill during the nine months ended September 30, 2012 resulted from the continued amortization of our intangible assets. This decrease was partly offset by the acquisition of a product right during the nine months ended September 30, 2012 and changes in foreign currency exchange rates.

NOTE 6. OTHER NONCURRENT ASSETS

Other noncurrent assets consisted of the following (*in thousands*):

	September 30, 2012	December 31, 2011
Investment in long-term product supply arrangements	\$ 10,750	\$ 12,091
Customer acquisition costs, net	22,488	21,075
Other assets	19,723	14,825
	<u>\$ 52,961</u>	<u>\$ 47,991</u>

NOTE 7. ACCRUED LIABILITIES

Accrued liabilities consisted of the following (*in thousands*):

	September 30, 2012	December 31, 2011
Accrued expenses	\$ 39,932	\$ 40,472
Accrued employee compensation and related expenses	50,801	51,373
Accrued taxes	14,890	17,654
Accrued customer programs	29,041	31,884
	<u>\$ 134,664</u>	<u>\$ 141,383</u>

NOTE 8. WARRANTY RESERVES

We provide a standard twelve month warranty on all instruments sold. We recognize the cost of instrument warranties in cost of product revenue at the time revenue is recognized based on the estimated cost to repair the instrument over its warranty period. Cost of product revenue reflects not only estimated warranty expense for instruments sold in the current period, but also any changes in estimated warranty expense for the portion of the aggregate installed base that is under warranty. Estimated warranty expense is based on a variety of inputs, including historical instrument performance in the customers' environment, historical costs incurred in servicing instruments and projected instrument reliability and service costs. Should actual service rates or costs differ from our estimates, revisions to the estimated warranty liability would be required. The liability for warranties is included in accrued liabilities in the accompanying condensed consolidated balance sheets.

The following is a summary of changes in accrued warranty reserves for the three and nine months ended September 30, 2012 and 2011 (*in thousands*):

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2012	2011	2012	2011
Balance, beginning of period	\$ 1,499	\$ 1,675	\$ 1,693	\$ 2,196
Provision for warranty expense	539	668	1,655	1,763
Change in estimate, balance beginning of period	7	(172)	(92)	(394)
Settlement of warranty liability	(613)	(604)	(1,824)	(1,998)
Balance, end of period	<u>\$ 1,432</u>	<u>\$ 1,567</u>	<u>\$ 1,432</u>	<u>\$ 1,567</u>

NOTE 9. REPURCHASES OF COMMON STOCK

The following is a summary of our open market common stock repurchases for the three and nine months ended September 30, 2012 and 2011 (*in thousands, except per share amounts*):

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2012	2011	2012	2011
Shares repurchased	386	886	1,038	2,183
Total cost of shares repurchased	\$ 36,146	\$ 67,597	\$ 91,152	\$ 166,016
Average cost per share	\$ 93.76	\$ 76.27	\$ 87.82	\$ 76.04

We primarily acquire shares by means of repurchases in the open market. However, we also acquire shares that are surrendered by employees in payment for the minimum required withholding taxes due on the vesting of restricted stock units and the settlement of deferred stock units. The number of shares acquired through employee surrenders during both the three months ended September 30, 2012 and 2011 was not material. We acquired 51,484 shares at a total cost of \$4.5 million in connection with such employee surrenders during the nine months ended September 30, 2012 compared to 54,940 shares at a total cost of \$4.3 million during the nine months ended September 30, 2011.

We issue shares of treasury stock upon the vesting of certain restricted stock units and upon the exercise of certain stock options. The number of shares of treasury stock issued during both the nine months ended September 30, 2012 and 2011 was not material.

NOTE 10. INCOME TAXES

Our effective income tax rates were 31.4% and 31.7% for the three and nine months ended September 30, 2012 compared to 30.8% and 31.1% for the three and nine months ended September 30, 2011, respectively. The increase in our effective income tax rate was due primarily to federal research and development tax incentives that were not available during the three and nine months ended September 30, 2012 but were available during the same periods of the prior year.

NOTE 11. ACCUMULATED OTHER COMPREHENSIVE INCOME

Accumulated other comprehensive income consisted of the following (*in thousands*):

	For the Nine Months Ended September 30, 2012		
	Unrealized loss on investments, net of tax	Unrealized gain (loss) on derivatives instruments, net of tax	Cumulative translation adjustment
Beginning balance	\$ (287)	\$ 3,206	\$ 12,524
Current-period other comprehensive income (loss)	111	(5,523)	4,548
Ending balance	<u>\$ (176)</u>	<u>\$ (2,317)</u>	<u>\$ 17,072</u>

NOTE 12. EARNINGS PER SHARE

Basic earnings per share is computed by dividing net income attributable to IDEXX Laboratories, Inc. stockholders by the weighted average number of shares of common stock and vested deferred stock units outstanding during the year. The computation of diluted earnings per share is similar to the computation of basic earnings per share, except that the denominator is increased for the assumed exercise of dilutive options and assumed issuance of unvested restricted stock units and unvested deferred stock units using the treasury stock method unless the effect is anti-dilutive. Vested deferred stock units outstanding are included in shares outstanding for basic and diluted earnings per share because the associated shares of our common stock are issuable for no cash consideration, the number of shares of our common stock to be issued is fixed and issuance is not contingent. See Note 4 to the consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2011 for additional information regarding deferred stock units.

The following is a reconciliation of shares outstanding for basic and diluted earnings per share for the three and nine months ended September 30, 2012 and 2011 (*in thousands*):

	For the Three Months Ended		For the Nine Months Ended	
	September 30,		September 30,	
	2012	2011	2012	2011
Shares outstanding for basic earnings per share:	54,938	56,699	55,074	57,141
Shares outstanding for diluted earnings per share:				
Shares outstanding for basic earnings per share	54,938	56,699	55,074	57,141
Dilutive effect of share-based payment awards	1,150	1,308	1,196	1,495
	<u>56,088</u>	<u>58,007</u>	<u>56,270</u>	<u>58,636</u>

Certain options to acquire shares have been excluded from the calculation of shares outstanding for diluted earnings per share because they were anti-dilutive. The following table presents information concerning those anti-dilutive options for the three and nine months ended September 30, 2012 and 2011 (*in thousands*):

	For the Three Months Ended		For the Nine Months Ended	
	September 30,		September 30,	
	2012	2011	2012	2011
Weighted average number of shares underlying anti-dilutive options	380	665	713	566

NOTE 13. COMMITMENTS, CONTINGENCIES AND GUARANTEES

As discussed in our Annual Report on Form 10-K for the year ended December 31, 2011, in January 2010, we received a letter from the U.S. Federal Trade Commission (“FTC”), stating that it was conducting an investigation to determine whether IDEXX or others have engaged in, or are engaging in, unfair methods of competition in violation of Section 5 of the Federal Trade Commission Act (“FTC Act”), through pricing or marketing policies for companion animal veterinary products and services, including but not limited to exclusive dealing or tying arrangements with distributors or end-users of those products or services. The letter requested that we preserve all materials potentially relevant to this investigation. The letter stated that the FTC had not concluded that IDEXX or anyone else had violated Section 5 of the FTC Act. We received subpoenas from the FTC on April 15, 2010 and August 8, 2011, requesting that we provide the FTC with documents and information relevant to this investigation and we have cooperated fully with the FTC in its investigation.

Based on discussions with the FTC staff, we determined that their primary concern was that our relationships with the three largest U.S. distributors of companion animal products, referred to as the national distributors, limited access by our competitors to companion animal veterinary practices. We also determined that their concerns could be addressed if one of these three national distributors was free to carry competitive products. In response to these concerns, in August 2012 we announced our intention to seek to modify our agreement with one of these distributors to eliminate our competitive products policy, which permits us to discontinue sale of our products in a particular product category to the distributor if the distributor sells competitive products in that product category. On September 28, 2012, we entered into such a modified agreement with MWI Veterinary Supply, Inc. (“MWI”) that will become effective January 1, 2013. Under this modified agreement, MWI will be permitted to carry any competitive products without restriction or potential negative consequence. This agreement was reviewed by the FTC staff

prior to its execution by IDEXX and MWI to ensure that the agreement met the staff's objective of ensuring that one national distributor had the right to carry competitive products without limitation.

We are currently negotiating the terms of a consent order with the FTC staff that we expect will require, among other things, that at least one national distributor will not be exclusive to IDEXX for the duration of the consent order, which is expected to be ten years. If successfully negotiated with the staff, the consent order would need to be approved by the FTC Commissioners prior to effectiveness. We cannot provide any assurance that we will be able to successfully negotiate the consent order with the staff, or that the FTC Commissioners will approve the consent order.

If a consent order is not approved by the FTC Commissioners, we cannot provide any assurances that the FTC will not determine to file a complaint against IDEXX in the administrative law court within the FTC at some point in the future, or that we would be successful defending ourselves in such a proceeding. Were we to be unsuccessful in defending this proceeding and any applicable appeals, we could be subject to restrictions on certain of our marketing and sales practices, including on the terms included in our agreements with certain of our U.S. distributors. While we cannot be certain about what prospective remedies would be sought by the FTC in any such proceeding, we believe that any required changes in our marketing or sales practices would not have a material adverse effect on our financial statements.

We continue to believe that our marketing and sales practices for companion animal veterinary products and services do not violate applicable antitrust laws. We have chosen to enter into the modified agreement with MWI and negotiate a consent order with the FTC staff because we believe this course will help us avoid long and costly litigation, and that our business will not be materially adversely affected.

Other significant commitments, contingencies and guarantees at September 30, 2012 are consistent with those discussed in Note 14 to the consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2011.

NOTE 14. SEGMENT REPORTING

We operate primarily through three business segments: diagnostic and information technology-based products and services for the veterinary market, which we refer to as the Companion Animal Group ("CAG"), water quality products ("Water") and products for livestock and poultry health, which we refer to as Livestock and Poultry Diagnostics ("LPD"). We also operate two smaller operating segments that comprise products for milk quality and safety ("Dairy") and products for the human point-of-care medical diagnostics market ("OPTI Medical"). Financial information about our Dairy and OPTI Medical operating segments is combined and presented with our remaining pharmaceutical product line and our out-licensing arrangements in an "Other" category because they do not meet the quantitative or qualitative thresholds for reportable segments.

The following is a summary of segment performance for the three and nine months ended September 30, 2012 and 2011 (in thousands):

	For the Three Months Ended September 30,					Unallocated Amounts	Consolidated Total
	CAG	Water	LPD	Other			
2012							
Revenue	\$ 262,357	\$ 22,223	\$ 18,911	\$ 11,984	\$ -	\$ 315,475	
Income (loss) from operations	\$ 50,651	\$ 10,128	\$ 3,504	\$ (119)	\$ (1,252)	\$ 62,912	
Interest expense, net						(400)	
Income before provision for income taxes						62,512	
Provision for income taxes						19,639	
Net income						42,873	
Less: Net income attributable to noncontrolling interest						20	
Net income attributable to IDEXX Laboratories, Inc. stockholders						\$ 42,853	

2011						
Revenue	\$ 248,074	\$ 21,648	\$ 20,675	\$ 10,557	\$ -	\$ 300,954
Income (loss) from operations	\$ 44,296	\$ 9,979	\$ 3,648	\$ 34	\$ (1,861)	\$ 56,096
Interest expense, net						(478)
Income before provision for income taxes						55,618
Provision for income taxes						17,122
Net income						38,496
Less: Net loss attributable to noncontrolling interest						(11)
Net income attributable to IDEXX Laboratories, Inc. stockholders						\$ 38,507

	For the Nine Months Ended September 30,					Consolidated Total
	CAG	Water	LPD	Other	Unallocated Amounts	
2012						
Revenue	\$ 808,724	\$ 63,788	\$ 64,153	\$ 37,135	\$ -	\$ 973,800
Income (loss) from operations	\$ 157,337	\$ 28,619	\$ 14,580	\$ 669	\$ (2,069)	\$ 199,136
Interest expense, net						(1,603)
Income before provision for income taxes						197,533
Provision for income taxes						62,606
Net income						134,927
Less: Net income attributable to noncontrolling interest						14
Net income attributable to IDEXX Laboratories, Inc. stockholders						\$ 134,913

2011						
Revenue	\$ 748,397	\$ 62,123	\$ 69,981	\$ 30,987	\$ -	\$ 911,488
Income (loss) from operations	\$ 145,137	\$ 25,327	\$ 17,974	\$ (207)	\$ (7,305)	\$ 180,926
Interest expense, net						(1,200)
Income before provision for income taxes						179,726
Provision for income taxes						55,970
Net income						123,756
Less: Net loss attributable to noncontrolling interest						(20)
Net income attributable to IDEXX Laboratories, Inc. stockholders						\$ 123,776

The following is a summary of revenue by product and service category for the three and nine months ended September 30, 2012 and 2011 (*in thousands*):

	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2012	2011	2012	2011
CAG segment revenue:				
Instruments and consumables	\$ 101,325	\$ 99,719	\$ 307,565	\$ 292,209
Rapid assay products	39,315	36,073	128,556	118,883
Reference laboratory diagnostic and consulting services	101,449	94,027	309,696	282,242
Practice management and digital imaging systems	20,268	18,255	62,907	55,063
CAG segment revenue	262,357	248,074	808,724	748,397
Water segment revenue	22,223	21,648	63,788	62,123
LPD segment revenue	18,911	20,675	64,153	69,981
Other segment revenue	11,984	10,557	37,135	30,987
Total revenue	\$ 315,475	\$ 300,954	\$ 973,800	\$ 911,488

NOTE 15. FAIR VALUE MEASUREMENTS

U.S. GAAP defines fair value as the price that would be received to sell an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. U.S. GAAP requires an entity to maximize the use of observable inputs, where available, and minimize the use of unobservable inputs when measuring fair value.

The Company has certain financial assets and liabilities that are measured at fair value on a recurring basis, certain nonfinancial assets and liabilities that may be measured at fair value on a nonrecurring basis and certain financial assets and liabilities that are not measured at fair value in our condensed consolidated balance sheets but for which we disclose the fair value. The fair value disclosures of these assets and liabilities are based on a three-level hierarchy, which is defined as follows:

- Level 1** Quoted prices (unadjusted) in active markets for identical assets or liabilities that the entity can access at the measurement date.
- Level 2** Observable inputs other than Level 1 prices, such as quoted prices for similar assets or liabilities; quoted prices in markets that are not active; or other inputs that are observable or can be corroborated by observable market data for substantially the full term of the assets or liabilities.
- Level 3** Unobservable inputs that are supported by little or no market activity and that are significant to the fair value of the assets or liabilities.

Assets and liabilities measured at fair value are classified in their entirety based on the lowest level of input that is significant to the fair value measurement. Our assessment of the significance of a particular input to the fair value measurement in its entirety requires judgment and considers factors specific to the asset or liability. We did not have any significant nonfinancial assets or nonfinancial liabilities which required remeasurement during the nine months ended September 30, 2012. We did not have any transfers between Level 1 and Level 2 or transfers in or out of Level 3 of the fair value hierarchy during the nine months ended September 30, 2012.

Our foreign currency exchange contracts and interest rate swap agreements are measured at fair value on a recurring basis in our accompanying condensed consolidated balance sheets. We measure the fair value of our foreign currency exchange contracts classified as derivative instruments using an income approach, based on prevailing market forward rates less the contract rate multiplied by the notional amount. The product of this calculation is then adjusted for counterparty risk. We measure the fair value of our interest rate swaps classified as derivative instruments using an income approach, utilizing a discounted cash flow analysis based on the terms of the contract and the interest rate curve adjusted for counterparty risk.

The amount outstanding under our unsecured revolving credit facility ("Credit Facility"), notes receivable and long-term debt are not measured at fair value in our accompanying condensed consolidated balance sheets. We determine the fair value of the amount outstanding under our Credit Facility, notes receivable and long-term debt using an income approach, utilizing a discounted cash flow analysis based on current market interest rates for debt issues with similar remaining years to maturity, adjusted for applicable credit risk. Our Credit Facility and long-term debt are valued using level 2 inputs, while our notes receivable, representing a strategic investment in a privately held company with a carrying value of \$4.5 million as of September 30, 2012, is valued using level 3 inputs. The results of these calculations yield fair values that approximate carrying values.

The following tables set forth our assets and liabilities that were measured at fair value on a recurring basis at September 30, 2012 and at December 31, 2011 by level within the fair value hierarchy (*in thousands*):

<u>As of September 30, 2012</u>	<u>Quoted Prices in Active Markets for Identical Assets (Level 1)</u>	<u>Significant Other Observable Inputs (Level 2)</u>	<u>Significant Unobservable Inputs (Level 3)</u>	<u>Balance at September 30, 2012</u>
Assets				
Money market funds ⁽¹⁾	\$ 134,536	\$ -	\$ -	\$ 134,536
Equity mutual funds ⁽²⁾	2,249	-	-	2,249
Foreign currency exchange contracts ⁽³⁾	-	2,585	-	2,585
Liabilities				
Foreign currency exchange contracts ⁽³⁾	-	2,338	-	2,338
Deferred compensation ⁽⁴⁾	2,249	-	-	2,249
Interest rate swaps ⁽⁵⁾	-	2,798	-	2,798

<u>As of December 31, 2011</u>	<u>Quoted Prices in Active Markets for Identical Assets (Level 1)</u>	<u>Significant Other Observable Inputs (Level 2)</u>	<u>Significant Unobservable Inputs (Level 3)</u>	<u>Balance at December 31, 2011</u>
Assets				
Money market funds ⁽¹⁾	\$ 88,525	\$ -	\$ -	\$ 88,525
Equity mutual funds ⁽²⁾	2,056	-	-	2,056
Foreign currency exchange contracts ⁽³⁾	-	6,841	-	6,841
Liabilities				
Foreign currency exchange contracts ⁽³⁾	-	1,753	-	1,753
Deferred compensation ⁽⁴⁾	2,056	-	-	2,056
Interest rate swaps ⁽⁵⁾	-	1,417	-	1,417

- (1) Money market funds are included within cash and cash equivalents. The remaining balance of cash and cash equivalents as of September 30, 2012 and December 31, 2011 was demand deposits.
- (2) Equity mutual funds relate to a deferred compensation plan that was assumed as part of a previous business combination. This amount is included within other long-term assets, net. See number (4) below for a discussion of the related deferred compensation liability.
- (3) Foreign currency exchange contracts are included within other current assets; other long-term assets, net; accrued liabilities; or other long-term liabilities depending on the gain (loss) position and anticipated settlement date.
- (4) Deferred compensation plans are included within other long-term liabilities. The fair value of our deferred compensation plan is indexed to the performance of the underlying equity mutual funds discussed in number (2) above.
- (5) Interest rate swaps are included within accrued liabilities.

The estimated fair value of certain financial instruments, including cash and cash equivalents, accounts receivable and accounts payable, approximate carrying value due to their short maturity.

NOTE 16. DERIVATIVE INSTRUMENTS AND HEDGING

Disclosure within this footnote is presented to provide transparency about how and why we use derivative instruments, how the instruments and related hedged items are accounted for, and how the instruments and related hedged items affect our financial position, results of operations and cash flows.

We are exposed to certain risks related to our ongoing business operations. The primary risks that we manage by using derivative instruments are foreign currency exchange risk and interest rate risk. Our subsidiaries enter into foreign currency exchange contracts to manage the exchange risk associated with their forecasted intercompany inventory purchases and sales for the next year. From time to time, we may also enter into foreign currency exchange contracts to minimize the impact of foreign currency fluctuations associated with specific, significant transactions. We enter into interest rate swaps to minimize the impact of interest rate fluctuations associated with our variable-rate debt.

The primary purpose of our foreign currency hedging activities is to protect against the volatility associated with foreign currency transactions. We also utilize natural hedges to mitigate our transaction and commitment exposures. Our corporate policy prescribes the range of allowable hedging activity. We enter into foreign currency exchange contracts with large multinational financial institutions and we do not hold or engage in transactions involving derivative instruments for purposes other than risk management.

We recognize all derivative instruments, including our foreign currency exchange contracts and interest rate swap agreements, on the balance sheet at fair value at the balance sheet date. Derivative instruments that do not qualify for hedge accounting treatment must be recorded at fair value through earnings. To qualify for hedge accounting treatment, cash flow hedges must be highly effective in offsetting changes to expected future cash flows on hedged transactions. If a derivative instrument qualifies for hedge accounting, changes in the fair value of the derivative instrument from the effective portion of the hedge are deferred in other comprehensive income ("OCI"), net of tax, and reclassified into earnings in the same period or periods during which the hedged transaction affects earnings. We immediately record in earnings the extent to which a hedge instrument is not effective in achieving offsetting changes in fair value. We de-designate derivative instruments from hedge accounting when the likelihood of the hedged transaction occurring becomes less than probable. For de-designated instruments, the gain or loss from the time of de-designation through maturity of the instrument is recognized in earnings. Any gain or loss in OCI at the time of de-designation is reclassified into earnings in the same period or periods during which the hedged transaction affects earnings. We enter into master netting arrangements with the counterparties to our derivative transactions which permit outstanding receivables and payables with the counterparties to our derivative transactions to be offset in the event of default. We present our derivative

assets and liabilities on a gross basis. All cash flows related to our foreign currency exchange contracts and interest rate swaps are classified as operating cash flows, which is consistent with the cash flow treatment of the underlying items being hedged.

Cash Flow Hedges

We have designated our foreign currency exchange contracts and variable-to-fixed interest rate swaps as cash flow hedges as these derivative instruments mitigate the exposure to variability in the cash flows of forecasted transactions attributable to foreign currency exchange and interest rates. Unless noted otherwise, we have also designated our derivative instruments as qualifying for hedge accounting treatment.

We did not de-designate any instruments from hedge accounting treatment during the three and nine months ended September 30, 2012 or 2011. Gains or losses related to hedge ineffectiveness recognized in earnings during the three and nine months ended September 30, 2012 and 2011 were not material. At September 30, 2012, the estimated amount of net losses that are expected to be reclassified out of accumulated OCI and into earnings within the next 12 months is \$2.2 million if exchange and interest rates do not fluctuate from the levels at September 30, 2012.

We enter into foreign currency exchange contracts for amounts that are less than the full value of forecasted intercompany inventory purchases and sales. Our hedging strategy related to intercompany inventory purchases and sales is to employ the full amount of our hedges for the succeeding year at the conclusion of our budgeting process for that year, which is complete by the end of the preceding year. We primarily utilize foreign currency exchange contracts with durations of less than 24 months. Quarterly, we enter into contracts to hedge incremental portions of anticipated foreign currency transactions for the current and following year. As a result, our risk with respect to foreign currency exchange rate fluctuations and the notional value of foreign currency exchange contracts may vary throughout the year.

We have entered into forward fixed interest rate swap agreements to manage the economic effect of variable interest obligations on amounts borrowed under the terms of our Credit Facility. Beginning on March 30, 2012, the variable interest rate associated with \$40 million of borrowings outstanding under the Credit Facility became effectively fixed at 1.36% plus the range of applicable interest rates ("Credit Spread") through September 30, 2016. Beginning on March 28, 2013, the variable interest rate associated with an additional \$40 million of borrowings outstanding under the Credit Facility will become effectively fixed at 1.64% plus the Credit Spread through September 30, 2016. Also on March 30, 2012, two of our forward fixed interest rate swap agreements expired. Under these agreements, the variable interest rate associated with \$80 million of borrowings outstanding under the Credit Facility had been effectively fixed at 2% plus the Credit Spread above the London interbank offered rate.

The U.S. dollar is the currency purchased or sold in all of our foreign currency exchange contracts. The notional amount of foreign currency exchange contracts to hedge forecasted intercompany inventory purchases and sales consisted of the following (*in thousands*):

Currency Sold	U.S. Dollar Equivalent	
	September 30, 2012	December 31, 2011
Euro	\$ 70,716	\$ 68,275
British pound	26,505	25,260
Canadian dollar	24,279	19,902
Australian dollar	14,274	12,417
Japanese yen	18,335	18,005
	<u>\$ 154,109</u>	<u>\$ 143,859</u>

Currency Purchased	U.S. Dollar Equivalent	
	September 30, 2012	December 31, 2011
Swiss franc	\$ 15,143	\$ 17,909

The notional amount of forward fixed interest rate swap agreements to manage variable interest obligations consisted of the following (*in thousands*):

	<u>U.S. Dollar Equivalent</u>	
	<u>September 30, 2012</u>	<u>December 31, 2011</u>
Interest rate swaps commencing March 31, 2010 and expiring March 30, 2012	\$ -	\$ 80,000
Interest rate swap commencing March 30, 2012 and expiring June 30, 2016	\$ 40,000	\$ 40,000
Interest rate swap commencing March 28, 2013 and expiring June 30, 2016	\$ 40,000	\$ 40,000

The fair values of derivative instruments and their respective classification in the condensed consolidated balance sheet consisted of the following (*in thousands*):

	<u>Balance Sheet Classification</u>	<u>Asset Derivatives</u>	
		<u>September 30, 2012</u>	<u>December 31, 2011</u>
Derivatives designated as hedging instruments			
Foreign currency exchange contracts	Other current assets	\$ 2,332	\$ 6,841
Foreign currency exchange contracts	Other long-term assets, net	253	-
Total derivative instruments		<u>\$ 2,585</u>	<u>\$ 6,841</u>

	<u>Balance Sheet Classification</u>	<u>Liability Derivatives</u>	
		<u>September 30, 2012</u>	<u>December 31, 2011</u>
Derivatives designated as hedging instruments			
Foreign currency exchange contracts	Accrued expenses	\$ 1,900	\$ 1,753
Foreign currency exchange contracts	Other long-term liabilities	438	-
Interest rate swaps	Accrued expenses	2,798	1,417
Total derivative instruments		<u>\$ 5,136</u>	<u>\$ 3,170</u>

The effect of derivative instruments designated as cash flow hedges on the condensed consolidated balance sheet for the three and nine months ended September 30, 2012 and 2011 consisted of the following (*in thousands*):

<u>Derivative instruments</u>	<u>Gain (Loss) Recognized in OCI on Derivative Instruments (Effective Portion)</u>			
	<u>For the Three Months Ended</u>		<u>For the Nine Months Ended</u>	
	<u>September 30,</u>	<u>September 30,</u>	<u>September 30,</u>	<u>September 30,</u>
	<u>2012</u>	<u>2011</u>	<u>2012</u>	<u>2011</u>
Foreign currency exchange contracts, net of tax	\$ (3,227)	\$ 5,324	\$ (4,655)	\$ 4,348
Interest rate swaps, net of tax	(329)	(57)	(868)	296
Total derivative instruments, net of tax	<u>\$ (3,556)</u>	<u>\$ 5,267</u>	<u>\$ (5,523)</u>	<u>\$ 4,644</u>

The effect of derivative instruments designated as cash flow hedges on the condensed consolidated statement of operations for the three and nine months ended September 30, 2012 and 2011 consisted of the following (*in thousands*):

<u>Derivative instruments</u>	<u>Classification of Gain (Loss) Reclassified from OCI into Income (Effective Portion)</u>	<u>Gain (Loss) Reclassified from Accumulated OCI into Income (Effective Portion)</u>			
		<u>For the Three Months Ended</u>		<u>For the Nine Months Ended</u>	
		<u>September 30,</u>	<u>September 30,</u>	<u>September 30,</u>	<u>September 30,</u>
		<u>2012</u>	<u>2011</u>	<u>2012</u>	<u>2011</u>
Foreign currency exchange contracts	Cost of revenue	\$ 1,379	\$ (1,385)	\$ 5,202	\$ (4,962)
Interest rate swaps	Interest expense	(112)	(371)	(574)	(1,070)
Total derivative instruments		<u>\$ 1,267</u>	<u>\$ (1,756)</u>	<u>\$ 4,628</u>	<u>\$ (6,032)</u>

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

This Quarterly Report on Form 10-Q contains statements which, to the extent they are not statements of historical fact, constitute "forward-looking statements." Such forward-looking statements about our business and expectations within the meaning of the Private Securities Litigation Reform Act of 1995, Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, include statements relating to future revenue growth rates, earnings and other measures of financial performance, the effect of economic conditions on our business performance; demand for our products; realizability of assets; future cash flow and uses of cash; future repurchases of common stock; future levels of indebtedness and capital spending; interest expense; warranty expense; share-based compensation expense; and competition. Forward-looking statements can be identified by the use of words such as "expects," "may," "anticipates," "intends," "would," "will," "plans," "believes," "estimates," "should," and similar words and expressions. These forward-looking statements are intended to provide our current expectations or forecasts of future events; are based on current estimates, projections, beliefs, and assumptions; and are not guarantees of future performance. Actual events or results may differ materially from those described in the forward-looking statements. These forward-looking statements involve a number of risks and uncertainties as more fully described under the heading "Part II, Item 1A. Risk Factors" in this Quarterly Report on Form 10-Q. The risks and uncertainties discussed herein do not reflect the potential impact of any mergers, acquisitions or dispositions. In addition, any forward-looking statements represent our estimates only as of the day this Quarterly Report was first filed with the Securities and Exchange Commission ("SEC") and should not be relied upon as representing our estimates as of any subsequent date. From time to time, oral or written forward-looking statements may also be included in other materials released to the public. While we may elect to update forward-looking statements at some point in the future, we specifically disclaim any obligation to do so, even if our estimates or expectations change.

■ Business Overview and Trends

Operating segments. We operate primarily through three business segments: diagnostic and information technology-based products and services for the veterinary market, which we refer to as the Companion Animal Group ("CAG"), water quality products ("Water") and products for livestock and poultry health, which we refer to as Livestock and Poultry Diagnostics ("LPD"). We also operate two smaller operating segments that comprise products for testing milk quality and safety ("Dairy") and products for the human point-of-care medical diagnostics market ("OPTI Medical"). Financial information about our Dairy and OPTI Medical operating segments is combined and presented with our remaining pharmaceutical product line and our out-licensing arrangements in an "Other" category because they do not meet the quantitative or qualitative thresholds for reportable segments. See Note 14 to the condensed consolidated financial statements included in this Quarterly Report on Form 10-Q for financial information about our segments and the section entitled "Description of Business by Segment" under the heading "Item 1. Business" in our Annual Report on Form 10-K for the year ended December 31, 2011 for additional description of our segments.

CAG develops, designs, manufactures and distributes products and performs services, primarily related to diagnostics and information management, for veterinarians. Water develops, designs, manufactures and distributes a range of products used in the detection of various microbiological parameters in water. LPD develops, designs, manufactures and distributes diagnostic tests and related instrumentation that are used to detect a wide range of diseases and to monitor health status in livestock and poultry. Dairy develops, designs, manufactures and distributes products to detect contaminants in milk. OPTI Medical develops, designs, manufactures and distributes point-of-care electrolyte and blood gas analyzers and related consumable products for the human medical diagnostics market. Further, OPTI Medical manufactures our VetStat[®] Electrolyte and Blood Gas Analyzer, a component of our Catalyst Dx[®] Analyzer and electrolyte consumables used with our Catalyst Dx[®] Analyzer.

Effects of Certain Factors on Results of Operations

Distributor Purchasing and Inventories. The instrument consumables and rapid assay products in our CAG segment are sold in the U.S. and certain other geographies by third party distributors, who purchase products from us and sell them to veterinary practices, which are the end users. As a result, distributor purchasing dynamics have an impact on our reported sales of these products. Distributor purchasing dynamics may be affected by many factors and in a given period may not be directly related to underlying end-user demand for our products. Consequently, reported results may reflect fluctuations in inventory levels held at distributors and not necessarily reflect changes in underlying end-user demand. Therefore, we believe it is important to track distributor sales to end users and to distinguish between the impact of end-user demand and the impact of distributor purchasing dynamics on reported revenue.

Where growth rates are affected by changes in end-user demand, we refer to this as the impact of practice-level sales on growth. Where growth rates are affected by distributor purchasing dynamics, we refer to this as the impact of changes in distributors' inventories on growth. If during the current year, distributors' inventories grew by less than those inventories grew in the comparable period of the prior year, then changes in distributors' inventories have an unfavorable impact on our reported sales growth in the current period. Conversely, if during the current year, distributors' inventories grew by more than those inventories grew in the comparable period of the prior year, then changes in distributors' inventories have a favorable impact on our reported sales growth in the current period.

At the end of a quarter, we believe that our U.S. CAG distributors typically hold inventory equivalent to approximately four weeks of the anticipated end-user demand for instrument consumables and rapid assay products.

Currency Impact. For both the three and nine months ended September 30, 2012, approximately 25% of our revenue was derived from products manufactured in the U.S. and sold internationally in local currencies compared to 25% and 26% for the three and nine months ended September 30, 2011, respectively. Strengthening of the rate of exchange for the U.S. dollar relative to other currencies has a negative impact on our revenues derived in currencies other than the U.S. dollar and on profits of products manufactured in the U.S. and sold internationally, and a weakening of the U.S. dollar has the opposite effect. Similarly, to the extent that the U.S. dollar is stronger in current or future periods relative to the exchange rates in effect in the corresponding prior periods, our growth rate will be negatively affected. The impact of foreign currency denominated operating expenses and foreign currency denominated supply contracts partly offset this exposure.

The impact on revenue resulting from changes in foreign currency exchange rates is not a measure defined by generally accepted accounting principles in the United States of America ("U.S. GAAP"), otherwise referred to herein as a non-U.S. GAAP measure. We calculate the impact on revenue resulting from changes in foreign currency exchange rates by applying the difference between the weighted average exchange rates during the current year period and the comparable previous year period to foreign currency denominated revenues for the current year period.

During the three months ended September 30, 2012, as compared to the three months ended September 30, 2011, changes in foreign currency exchange rates decreased total company revenue by approximately \$7.6 million, due primarily to the strengthening of the U.S. dollar against the Euro.

During the nine months ended September 30, 2012, as compared to the nine months ended September 30, 2011, changes in foreign currency exchange rates decreased total company revenue by approximately \$17.4 million, due primarily to the strengthening of the U.S. dollar against the Euro.

Effects of Economic Conditions. Demand for many of our products and services has been negatively affected by economic conditions since mid-2008. In our CAG segment, we believe that low economic growth and relatively high unemployment have led to negative or cautious consumer sentiment, which has affected the number of patient visits with their respective owners to veterinary clinics. Based on data provided by a subset of our U.S. customers that use our practice management systems, we observed a slight improvement in the growth of patient visits beginning in the fourth quarter of 2011 and further improvement during 2012 over the previous year periods, although the rate of improvement has not been steady. We believe that this data, though limited, provides a fair and meaningful representation of patient visit activity in the U.S. that may be indicative of a gradual but inconsistent improvement in the U.S. economic environment. In contrast, economic conditions in certain European countries remain challenging, which continues to negatively impact our CAG segment in particular.

We believe that the overall trend in patient visits since the beginning of the economic downturn has had a slightly negative impact on the growth rate of sales of rapid assay tests, instrument consumables and reference laboratory diagnostic and consulting services. In addition, we believe the rate of growth of sales of our instruments and digital radiography systems, which are larger capital purchases for veterinarians, has also been affected by continued caution among veterinarians regarding economic conditions. Weaker economic conditions since mid-2008 have also caused our customers to remain sensitive to the pricing of our products and services resulting in lower revenue growth due to limited price increases for certain products.

We also believe that current economic conditions have affected purchasing decisions of our Water and LPD business customers. Lower water testing volumes have resulted from a decline in discretionary testing and a decline in mandated testing as a result of lower home and commercial construction. Fiscal difficulties in certain European countries have also reduced government funding for some water and livestock testing programs.

We believe that the diversity of our products and services and the geographic diversity of our markets have partially mitigated the effects of the economic environment and negative consumer sentiment on our revenue growth rates. Looking forward, we are cautiously optimistic that the improvements we began to see in the U.S. commencing in the fourth quarter of 2011 and continuing in 2012 are reflective of a gradual improvement in the macroeconomic environment that over time will further reduce these effects.

■ Critical Accounting Policies and Estimates

The discussion and analysis of our financial condition and results of operations is based upon our condensed consolidated financial statements, which have been prepared in accordance with U.S. GAAP. The preparation of these financial statements requires us to make estimates and judgments that affect the reported amounts of assets, liabilities, revenues and expenses, and related disclosure of contingent assets and liabilities. On an ongoing basis, we evaluate our estimates. We base our estimates on historical experience and on various assumptions that we believe to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Actual results may differ from these estimates. The significant accounting policies used in preparation of these condensed consolidated financial statements for the three and nine months ended September 30, 2012 are consistent with those discussed in Note 2 to the consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2011, except for our significant accounting policies related to revenue recognition. Effective January 1, 2012, revenue from substantially all U.S. distributors is recognized upon delivery to the distributor because title and risk of loss remains with IDEXX until the product is delivered. Prior to January 1, 2012, we recognized revenue at the time of shipment to U.S. distributors because title and risk of loss passed to the distributors on delivery to the common carrier. This change did not have a material impact on our financial statements. The critical accounting policies and the significant judgments and estimates used in the preparation of our condensed consolidated financial statements for the three and nine months ended September 30, 2012 are consistent with those discussed in our Annual Report on Form 10-K for the year ended December 31, 2011 in the section under the heading “Part 2, Item 7. Management’s Discussion and Analysis of Financial Condition and Results of Operations—Critical Accounting Policies and Estimates.”

■ Results of Operations

The following revenue analysis and discussion reports on organic revenue growth. Organic revenue growth should be considered in addition to, and not as a replacement for or as a superior measure to, revenues reported in accordance with U.S. GAAP, and may not be comparable to similarly titled measures reported by other companies. Management believes that reporting organic revenue growth provides useful information to investors by facilitating easier comparisons of our revenue performance with prior and future periods and to our peers. We exclude the effect of changes in foreign currency exchange rates because changes in foreign currency exchange rates are not under management’s control, are subject to volatility and can obscure underlying business trends. We exclude the effect of acquisitions because the nature, size and number of acquisitions can vary dramatically from period to period and therefore can also obscure underlying business trends.

Organic revenue growth and the percentage changes in revenue from currency and acquisitions are non-U.S. GAAP measures. See the subsection above titled “Business Overview and Trends” for a description of the calculation of the percentage change in revenue resulting from changes in foreign currency exchange rates. The percentage change in revenue resulting from acquisitions represents incremental revenues attributable to acquisitions that have occurred since the beginning of the prior year period.

Three Months Ended September 30, 2012 Compared to Three Months Ended September 30, 2011

Revenue

Total Company. The following table presents revenue by operating segment:

<u>Net Revenue</u> <i>(dollars in thousands)</i>	<u>For the Three</u> <u>Months Ended</u> <u>September 30,</u> <u>2012</u>	<u>For the Three</u> <u>Months Ended</u> <u>September 30,</u> <u>2011</u>	<u>Dollar</u> <u>Change</u>	<u>Percentage</u> <u>Change</u>	<u>Percentage</u> <u>Change from</u> <u>Currency</u>	<u>Percentage</u> <u>Change from</u> <u>Acquisitions</u>	<u>Organic</u> <u>Revenue Growth</u>
CAG	\$ 262,357	\$ 248,074	\$ 14,283	5.8 %	(2.3 %)	1.4 %	6.7 %
Water	22,223	21,648	575	2.7 %	(2.0 %)	-	4.7 %
LPD	18,911	20,675	(1,764)	(8.5 %)	(6.1 %)	-	(2.4 %)
Other	11,984	10,557	1,427	13.5 %	(1.6 %)	-	15.1 %
Total Company	<u>\$ 315,475</u>	<u>\$ 300,954</u>	<u>\$ 14,521</u>	4.8 %	(2.5 %)	1.1 %	6.2 %

Companion Animal Group. The following table presents revenue by product and service category for CAG:

Net Revenue <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	For the Three Months Ended September 30, 2011	Dollar Change	Percentage Change	Percentage Change from Currency	Percentage Change from Acquisitions	Organic Revenue Growth
Instruments and consumables	\$ 101,325	\$ 99,719	\$ 1,606	1.6 %	(2.6 %)	-	4.2 %
Rapid assay products	39,315	36,073	3,242	9.0 %	(1.1 %)	-	10.1 %
Reference laboratory diagnostic and consulting services	101,449	94,027	7,422	7.9 %	(3.0 %)	3.7 %	7.2 %
Practice management and digital imaging systems	20,268	18,255	2,013	11.0 %	(0.1 %)	-	11.1 %
Net CAG revenue	<u>\$ 262,357</u>	<u>\$ 248,074</u>	<u>\$ 14,283</u>	5.8 %	(2.3 %)	1.4 %	6.7 %

Instruments revenue was \$21.7 million and \$24.8 million for the three months ended September 30, 2012 and 2011, respectively. Consumables revenue was \$67.7 million and \$63.3 million for the three months ended September 30, 2012 and 2011, respectively. Instrument service and accessories revenue was \$11.6 million and \$11.2 million for the three months ended September 30, 2012 and 2011, respectively. The remaining sources of revenue are not significant to overall instruments and consumables revenue for both the three months ended September 30, 2012 and 2011. The decrease in instruments revenue was due primarily to lower sales of our Catalyst Dx[®] instrument compared to the three months ended September 30, 2011, which benefited significantly from the initial launch of a Catalyst Dx[®] instrument marketing program in North America. Consumables revenue growth was due primarily to higher sales volumes of consumables used with our Catalyst Dx[®] instrument, partly offset by lower sales of consumables used with our VetTest[®] chemistry instrument as customers continue to upgrade from our VetTest[®] instrument to our Catalyst Dx[®] instrument. Higher sales volumes of consumables used with our ProCyte Dx[®] instrument also contributed to the increase in consumables revenue. Service and accessories revenue growth was primarily a result of the increase in our active installed base of instruments. The impact of changes in distributors' inventory levels contributed 1% to instruments and consumables revenue growth.

The increase in rapid assay revenue was due primarily to higher sales of our canine combination test products and the favorable impact of changes in distributors' inventory levels, which contributed 4% to revenue growth. The higher sales of our canine combination test products was the result of increased sales volumes and higher average unit sales prices.

The increase in reference laboratory diagnostic and consulting services revenue resulted primarily from the impact of higher testing volumes and, to a lesser extent, price increases. Higher testing volumes were driven by the acquisition of new customers due, in part, to geographic expansion and our customer loyalty programs in which customers are provided incentives in exchange for agreements to purchase services in future periods.

The increase in practice management and digital imaging systems revenue resulted primarily from incremental revenue recognized under customer loyalty programs where revenue had been deferred at the time of system placements and in increase in service and support revenue.

Water. The increase in Water revenue resulted primarily from higher Quanti-Tray[®] and Colilert[®] product sales volumes due to new account acquisitions.

Livestock and Poultry Diagnostics. The decrease in LPD revenue resulted primarily from lower sales volumes of certain bovine tests in Europe due to declines in certain government programs.

Other. The increase in Other revenue was attributable primarily to higher sales volumes of our Dairy SNAP[®] tests used for the detection of antibiotic residue and the contaminant Aflatoxin M1 in milk. Higher sales of our OPTI Medical instruments also contributed to the increase in revenue.

Gross Profit

Total Company. The following table presents gross profit and gross profit percentages by operating segment:

Gross Profit <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	Percent of Revenue	For the Three Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
CAG	\$ 138,558	52.8 %	\$ 126,048	50.8 %	\$ 12,510	9.9 %
Water	14,489	65.2 %	14,317	66.1 %	172	1.2 %
LPD	12,630	66.8 %	13,666	66.1 %	(1,036)	(7.6 %)
Other	4,577	38.2 %	4,009	38.0 %	568	14.2 %
Unallocated Amounts	381	N/A	627	N/A	(246)	N/A
Total Company	<u>\$ 170,635</u>	54.1 %	<u>\$ 158,667</u>	52.7 %	<u>\$ 11,968</u>	7.5 %

Companion Animal Group. Gross profit for CAG increased due to higher sales and an increase in the gross profit percentage to 53% from 51%. The increase in gross profit percentage was due primarily to lower overall costs driven by higher volumes of consumables used in our VetLab[®] instruments, the favorable impact of currency and higher relative sales of higher margin products. These favorable impacts were partly offset by higher costs to support the continued expansion of our reference laboratories network. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency exchange rates was more than offset by hedging gains during the three months ended September 30, 2012 compared to hedging losses during the same period of the prior year.

Water. Gross profit for Water increased due to higher sales, partly offset by a decrease in the gross profit percentage to 65% from 66%. The decrease in the gross profit percentage was due primarily to higher relative sales of our Quanti-Tray[®] products that yield lower margins and higher raw materials costs. These unfavorable impacts were partly offset by the favorable impact of currency. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency was more than offset by hedging gains during the three months ended September 30, 2012 compared to hedging losses during the same period of the prior year.

Livestock and Poultry Diagnostics. Gross profit for LPD decreased due to lower sales, partly offset by an increase in the gross profit percentage to 67% from 66%. The increase in the gross profit percentage was due primarily to higher relative sales of products that yield higher margins and the favorable impact of currency. These favorable impacts were partly offset by higher overall manufacturing costs driven by lower production volumes and an increase in distribution costs. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency exchange rates was more than offset by hedging gains during the three months ended September 30, 2012 compared to hedging losses during the same period of the prior year.

Other. Gross profit for Other increased due to higher sales and a slight increase in the gross profit percentage. The increase in the gross profit percentage was due to lower overall manufacturing costs driven primarily by our OPTI Medical line of business. This favorable factor was partly offset by lower average unit sales prices also driven by our OPTI Medical line of business.

Unallocated Amounts. Gross profit for Unallocated Amounts decreased due primarily to an increase in personnel-related costs, partly offset by a decrease in certain manufacturing costs. With respect to personnel-related costs, we estimate certain personnel-related costs and allocate the estimated expenses to the operating segments. This allocation differs from actual expense and consequently yields a difference that is reported under the caption "Unallocated Amounts." The increase in certain personnel-related costs is due primarily to higher self-insured health care costs during the three months ended September 30, 2012 compared to the same period of the prior year. The manufacturing costs reported in our operating segments include our standard cost for products sold and any variances from standard cost for products purchased or manufactured within the period. We capitalize these variances for inventory on hand at the end of the period to record inventory in accordance with U.S. GAAP. We then record these variances as cost of product revenue as that inventory is sold. The impact to cost of product revenue resulting from this variance capitalization and subsequent recognition is reported within the caption "Unallocated Amounts." The decrease in certain manufacturing costs was due primarily to the recognition of previously favorable purchase price variances incurred in our LPD business.

Operating Expenses and Operating Income

Total Company. The following tables present operating expenses and operating income by operating segment:

Operating Expenses <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	Percent of Revenue	For the Three Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
CAG	\$ 87,907	33.5 %	\$ 81,752	33.0 %	\$ 6,155	7.5 %
Water	4,361	19.6 %	4,338	20.0 %	23	0.5 %
LPD	9,126	48.3 %	10,018	48.5 %	(892)	(8.9 %)
Other	4,696	39.2 %	3,975	37.7 %	721	18.1 %
Unallocated Amounts	1,633	N/A	2,488	N/A	(855)	(34.4 %)
Total Company	<u>\$ 107,723</u>	34.1 %	<u>\$ 102,571</u>	34.1 %	<u>\$ 5,152</u>	5.0 %

Operating Income <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	Percent of Revenue	For the Three Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
CAG	\$ 50,651	19.3 %	\$ 44,296	17.9 %	\$ 6,355	14.3 %
Water	10,128	45.6 %	9,979	46.1 %	149	1.5 %
LPD	3,504	18.5 %	3,648	17.6 %	(144)	(3.9 %)
Other	(119)	(1.0 %)	34	0.3 %	(153)	(450.0 %)
Unallocated Amounts	(1,252)	N/A	(1,861)	N/A	609	(32.7 %)
Total Company	<u>\$ 62,912</u>	19.9 %	<u>\$ 56,096</u>	18.6 %	<u>\$ 6,816</u>	12.2 %

Companion Animal Group. The following table presents CAG operating expenses by functional area:

Operating Expenses <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	Percent of Revenue	For the Three Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
Sales and marketing	\$ 45,110	17.2 %	\$ 43,327	17.5 %	\$ 1,783	4.1 %
General and administrative	28,965	11.0 %	25,724	10.4 %	3,241	12.6 %
Research and development	13,832	5.3 %	12,701	5.1 %	1,131	8.9 %
Total operating expenses	<u>\$ 87,907</u>	33.5 %	<u>\$ 81,752</u>	33.0 %	<u>\$ 6,155</u>	7.5 %

The increase in sales and marketing expense resulted primarily from higher personnel-related costs and increased spending on promotional activities, partly offset by the favorable impact from changes in foreign currency exchange rates. The increase in general and administrative expense resulted primarily from higher personnel-related costs. The increase in research and development expense resulted primarily from increased personnel-related costs and higher external consulting and development costs.

Water. The following table presents Water operating expenses by functional area:

Operating Expenses <i>(dollars in thousands)</i>	For the Three Months Ended September 30, 2012	Percent of Revenue	For the Three Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
Sales and marketing	\$ 2,233	10.0 %	\$ 2,114	9.8 %	\$ 119	5.6 %
General and administrative	1,552	7.0 %	1,595	7.4 %	(43)	(2.7 %)
Research and development	576	2.6 %	629	2.9 %	(53)	(8.4 %)
Total operating expenses	<u>\$ 4,361</u>	19.6 %	<u>\$ 4,338</u>	20.0 %	<u>\$ 23</u>	0.5 %

The increase in sales and marketing expense resulted primarily from higher personnel-related costs. General and administrative expense for the three months ended September 30, 2012 was generally consistent with the same period of the prior year. The decrease in research and development expense was due primarily to the timing of certain projects.

Livestock and Poultry Diagnostics. The following table presents LPD operating expenses by functional area:

<u>Operating Expenses</u> <i>(dollars in thousands)</i>	<u>For the Three Months Ended September 30, 2012</u>	<u>Percent of Revenue</u>	<u>For the Three Months Ended September 30, 2011</u>	<u>Percent of Revenue</u>	<u>Dollar Change</u>	<u>Percentage Change</u>
Sales and marketing	\$ 3,349	17.7 %	\$ 3,985	19.3 %	\$ (636)	(16.0 %)
General and administrative	2,829	15.0 %	2,905	14.1 %	(76)	(2.6 %)
Research and development	2,948	15.6 %	3,128	15.1 %	(180)	(5.8 %)
Total operating expenses	<u>\$ 9,126</u>	48.3 %	<u>\$ 10,018</u>	48.5 %	<u>\$ (892)</u>	(8.9 %)

The decrease in sales and marketing expense resulted primarily from the favorable impact of changes in foreign currency exchange rates and a decrease in personnel-related costs. The decrease in general and administrative expense resulted primarily from a decrease in intangible asset amortization and the favorable impact of changes in foreign currency exchange rates. The decrease in research and development expense was due primarily to the favorable impact of changes in foreign currency exchange rates and lower personnel-related costs.

Other. Operating expenses for Other increased \$0.7 million to \$4.7 million for the three months ended September 30, 2012 due primarily to higher personnel-related costs in our OPTI Medical line of business.

Unallocated Amounts. Operating expenses that are not allocated to our operating segments decreased \$0.9 million to \$1.6 million for the three months ended September 30, 2012 due primarily to lower legal and other professional fees incurred in connection with the U.S. Federal Trade Commission ("FTC") investigation, certain foreign exchange gains during the three months ended September 30, 2012 compared to losses during the same period of the prior year and a decrease in personnel-related costs related to unallocated corporate support function expenses. These favorable impacts were partly offset by the absence of a payment related to the sale of certain raw material inventory in connection with the restructuring of our pharmaceutical business in 2008 that was recognized during the three months ended September 30, 2011. We estimate corporate support function expenses and certain personnel-related costs and allocate the estimated expense to the operating segments. This allocation differs from the actual expense and consequently yields a difference that is reported under the caption "Unallocated Amounts." The FTC investigation is discussed in more detail under the heading "Part II. Item 1A. Risk Factors" in this Quarterly Report on Form 10-Q.

Interest Income and Interest Expense

Interest income was \$0.5 million for both the three months ended September 30, 2012 and 2011.

Interest expense was \$0.9 million for both the three months ended September 30, 2012 and 2011, as higher average balances outstanding on our unsecured revolving credit facility ("Credit Facility") were offset by lower effective interest rates.

Provision for Income Taxes

Our effective income tax rate was 31.4% for the three months ended September 30, 2012 compared to 30.8% for the three months ended September 30, 2011. The increase in our effective income tax rate was due primarily to federal research and development tax incentives that were not available during the three months ended September 30, 2012 but were available during the same period of the prior year.

Nine Months Ended September 30, 2012 Compared to Nine Months Ended September 30, 2011

The following revenue analysis and discussion reports on organic revenue growth. Organic revenue growth should be considered in addition to, and not as a replacement for or as a superior measure to, revenues reported in accordance with U.S. GAAP, and may not be comparable to similarly titled measures reported by other companies. Management believes that reporting organic revenue growth provides useful information to investors by facilitating easier comparisons of our revenue performance with prior and future periods and to our peers. We exclude the effect of changes in foreign currency exchange rates because changes in foreign currency exchange rates are not under management's control, are subject to volatility and can obscure underlying business trends. We exclude the effect of acquisitions because the nature, size and number of acquisitions can vary dramatically from period to period and therefore can also obscure underlying business trends.

Organic revenue growth and the percentage changes in revenue from currency and acquisitions are non-U.S. GAAP measures. See the subsection above titled “Business Overview and Trends” for a description of the calculation of the percentage change in revenue resulting from changes in foreign currency exchange rates. The percentage change in revenue resulting from acquisitions represents incremental revenues attributable to acquisitions that have occurred since the beginning of the prior year period.

Revenue

Total Company. The following table presents revenue by operating segment:

<i>Net Revenue (dollars in thousands)</i>	For the Nine Months Ended September 30, 2012	For the Nine Months Ended September 30, 2011	Dollar Change	Percentage Change	Percentage Change from Currency	Percentage Change from Acquisitions	Organic Revenue Growth
CAG	\$ 808,724	\$ 748,397	\$ 60,327	8.1 %	(1.7 %)	1.4 %	8.4 %
Water	63,788	62,123	1,665	2.7 %	(1.6 %)	-	4.3 %
LPD	64,153	69,981	(5,828)	(8.3 %)	(4.3 %)	-	(4.0 %)
Other	37,135	30,987	6,148	19.8 %	(0.9 %)	-	20.7 %
Total Company	<u>\$ 973,800</u>	<u>\$ 911,488</u>	<u>\$ 62,312</u>	6.8 %	(2.0 %)	1.2 %	7.6 %

Companion Animal Group. The following table presents revenue by product and service category for CAG:

<i>Net Revenue (dollars in thousands)</i>	For the Nine Months Ended September 30, 2012	For the Nine Months Ended September 30, 2011	Dollar Change	Percentage Change	Percentage Change from Currency	Percentage Change from Acquisitions	Organic Revenue Growth
Instruments and consumables	\$ 307,565	\$ 292,209	\$ 15,356	5.3 %	(2.0 %)	-	7.3 %
Rapid assay products	128,556	118,883	9,673	8.1 %	(0.8 %)	-	8.9 %
Reference laboratory diagnostic and consulting services	309,696	282,242	27,454	9.7 %	(2.2 %)	3.7 %	8.2 %
Practice management and digital imaging systems	62,907	55,063	7,844	14.2 %	(0.2 %)	-	14.4 %
Net CAG revenue	<u>\$ 808,724</u>	<u>\$ 748,397</u>	<u>\$ 60,327</u>	8.1 %	(1.7 %)	1.4 %	8.4 %

Instruments revenue was \$65.6 million and \$64.9 million for the nine months ended September 30, 2012 and 2011, respectively. Consumables revenue was \$206.4 million and \$193.5 million for the nine months ended September 30, 2012 and 2011, respectively. Instrument service and accessories revenue was \$34.4 million and \$32.8 million for the nine months ended September 30, 2012 and 2011, respectively. The remaining sources of revenue are not significant to overall instruments and consumables revenue. Instruments revenue growth was due primarily to increased sales of our ProCyte Dx[®] instrument. Consumables revenue growth was due primarily to higher sales volumes of consumables used with our Catalyst Dx[®] instrument, partly offset by lower sales of consumables used with our VetTest[®] chemistry instrument as customers continue to upgrade from our VetTest[®] instrument to our Catalyst Dx[®] instrument. Higher sales volumes of consumables used with our ProCyte Dx[®] instrument also contributed to the increase in consumables revenue, partly offset by lower sales of consumables used with our LaserCyte[®] instrument as customers continue to upgrade from our LaserCyte[®] instrument to our ProCyte Dx[®] instrument. Service and accessories revenue growth was primarily a result of the increase in our active installed base of instruments. The impact of changes in distributors’ inventory levels was not significant to reported instruments and consumables growth.

The increase in rapid assay revenue was due primarily to higher sales of our canine combination test products and, to a lesser extent, the favorable impact of changes in distributors’ inventory levels worldwide, which contributed 2% to revenue growth. Higher sales of our canine combination test products were driven primarily by higher average unit sales prices.

The increase in reference laboratory diagnostic and consulting services revenue resulted primarily from the impact of higher testing volumes. Higher testing volumes were driven by the acquisition of new customers due, in part, to geographic

expansion and our customer loyalty programs in which customers are provided incentives in exchange for agreements to purchase services in future periods.

The increase in practice management and digital imaging systems revenue resulted primarily from an increase in placements of our practice management systems, an increase in support and service revenue and incremental digital radiography revenue recognized under customer loyalty programs where revenue had been deferred at the time of system placement.

Water. The increase in Water revenue resulted primarily from higher Colilert® and Quanti-Tray® product sales volumes due to new account acquisitions.

Livestock and Poultry Diagnostics. The decrease in LPD revenue resulted primarily from lower sales of bovine tests, due in part to lower sales volumes of Bovine Spongiform Encephalopathy (“BSE” or “mad cow disease”) tests resulting from the changes in European Union BSE testing requirements. Effective July 1, 2011, the age at which healthy cattle to be slaughtered are required to be tested for BSE in the European Union was increased from 48 months to 72 months, which is reducing the population of cattle tested for this disease.

Other. The increase in Other revenue was attributable primarily to higher sales volumes of our Dairy SNAP® tests used for the detection of antibiotic residue and the contaminant Aflatoxin M1 in milk. Higher sales of our OPTI Medical instruments and related consumables also contributed to the increase in revenue.

Gross Profit

Total Company. The following table presents gross profit and gross profit percentages by operating segment:

<u>Gross Profit (dollars in thousands)</u>	<u>For the Nine Months Ended September 30, 2012</u>	<u>Percent of Revenue</u>	<u>For the Nine Months Ended September 30, 2011</u>	<u>Percent of Revenue</u>	<u>Dollar Change</u>	<u>Percentage Change</u>
CAG	\$ 425,867	52.7 %	\$ 387,734	51.8 %	\$ 38,133	9.8 %
Water	42,250	66.2 %	38,676	62.3 %	3,574	9.2 %
LPD	43,483	67.8 %	47,548	67.9 %	(4,065)	(8.5 %)
Other	14,469	39.0 %	12,493	40.3 %	1,976	15.8 %
Unallocated Amounts	4,029	N/A	1,174	N/A	2,855	N/A
Total Company	<u>\$ 530,098</u>	54.4 %	<u>\$ 487,625</u>	53.5 %	<u>\$ 42,473</u>	8.7 %

Companion Animal Group. Gross profit for CAG increased due to higher sales and an increase in the gross profit percentage to 53% from 52%. The increase in the gross profit percentage was due primarily to lower overall costs driven by higher volumes of consumables used in our VetLab® instruments, higher average unit sales prices of our canine combination rapid assay tests and the favorable impact of currency. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency exchange rates was more than offset by hedging gains during the nine months ended September 30, 2012 compared to hedging losses during the same period of the prior year. These favorable factors were partly offset by increased costs to support the continued expansion of our reference laboratories network.

Water. Gross profit for Water increased due to higher sales and an increase in the gross profit percentage to 66% from 62%. The increase in the gross profit percentage was due primarily to the timing of certain manufacturing costs during the nine months ended September 30, 2011 and the favorable impact of currency. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency exchange rates was more than offset by hedging gains during the nine months ended September 30, 2012 compared to hedging losses during the same period of the prior year.

Livestock and Poultry Diagnostics. Gross profit for LPD decreased due to lower sales and a slight decrease in the gross profit percentage. Higher overall manufacturing costs driven by lower production volumes were almost entirely offset by the favorable impact of currency and higher relative sales of higher margin products. The net effect of currency was positive because the net unfavorable impact of changes in foreign currency exchange rates was more than offset by hedging gains during the nine months ended September 30, 2012 compared to hedging losses during the same period of the prior year.

Other. Gross profit for Other increased due to higher sales, partly offset by a decrease in the gross profit percentage to 39% from 40%. The decrease in the gross profit percentage was due to lower average unit sales prices, driven by our OPTI Medical line of business, and higher distribution and freight costs in our Dairy and OPTI Medical lines of business. These

unfavorable factors were partly offset by lower overall manufacturing costs in our Dairy line of business driven by increased production volumes of our SNAP[®] tests.

Unallocated Amounts. Gross profit for Unallocated Amounts increased due primarily to a decrease in certain manufacturing costs. The manufacturing costs reported in our operating segments include our standard cost for products sold and any variances from standard cost for products purchased or manufactured within the period. We capitalize these variances for inventory on hand at the end of the period to record inventory in accordance with U.S. GAAP. We then record these variances as cost of product revenue as that inventory is sold. The impact to cost of product revenue resulting from this variance capitalization and subsequent recognition is reported within the caption “Unallocated Amounts.” The decrease in certain manufacturing costs is due primarily to the recognition of previously favorable production volume and purchase price variances incurred in our LPD business.

Operating Expenses and Operating Income

Total Company. The following tables present operating expenses and operating income by operating segment:

<i>Operating Expenses</i> <i>(dollars in thousands)</i>	For the Nine Months Ended September 30, 2012	Percent of Revenue	For the Nine Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
CAG	\$ 268,530	33.2 %	\$ 242,597	32.4 %	\$ 25,933	10.7 %
Water	13,631	21.4 %	13,349	21.5 %	282	2.1 %
LPD	28,903	45.1 %	29,574	42.3 %	(671)	(2.3 %)
Other	13,800	37.2 %	12,700	41.0 %	1,100	8.7 %
Unallocated Amounts	6,098	N/A	8,479	N/A	(2,381)	(28.1 %)
Total Company	\$ 330,962	34.0 %	\$ 306,699	33.7 %	\$ 24,263	7.9 %

<i>Operating Income</i> <i>(dollars in thousands)</i>	For the Nine Months Ended September 30, 2012	Percent of Revenue	For the Nine Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
CAG	\$ 157,337	19.5 %	\$ 145,137	19.4 %	\$ 12,200	8.4 %
Water	28,619	44.9 %	25,327	40.8 %	3,292	13.0 %
LPD	14,580	22.7 %	17,974	25.7 %	(3,394)	(18.9 %)
Other	669	1.8 %	(207)	(0.7 %)	876	(423.2 %)
Unallocated Amounts	(2,069)	N/A	(7,305)	N/A	5,236	(71.7 %)
Total Company	\$ 199,136	20.4 %	\$ 180,926	19.9 %	\$ 18,210	10.1 %

Companion Animal Group. The following table presents CAG operating expenses by functional area:

<i>Operating Expenses</i> <i>(dollars in thousands)</i>	For the Nine Months Ended September 30, 2012	Percent of Revenue	For the Nine Months Ended September 30, 2011	Percent of Revenue	Dollar Change	Percentage Change
Sales and marketing	\$ 140,713	17.4 %	\$ 129,489	17.3 %	\$ 11,224	8.7 %
General and administrative	86,235	10.7 %	76,941	10.3 %	9,294	12.1 %
Research and development	41,582	5.1 %	36,167	4.8 %	5,415	15.0 %
Total operating expenses	\$ 268,530	33.2 %	\$ 242,597	32.4 %	\$ 25,933	10.7 %

The increase in sales and marketing expense resulted primarily from higher personnel-related costs, partly offset by the favorable impact from changes in foreign currency exchange rates. The increase in general and administrative expense was due primarily to higher personnel-related costs, higher amortization expense of our intangible assets and an increase in costs attributable to investments in information technology. These unfavorable impacts were partly offset by the favorable impact of changes in foreign currency exchange rates. The increase in research and development expense resulted primarily from higher personnel-related costs and increased external consulting and development costs.

Water. The following table presents Water operating expenses by functional area:

<u>Operating Expenses</u> <i>(dollars in thousands)</i>	<u>For the Nine</u> <u>Months Ended</u> <u>September 30,</u> <u>2012</u>	<u>Percent of</u> <u>Revenue</u>	<u>For the Nine</u> <u>Months Ended</u> <u>September 30,</u> <u>2011</u>	<u>Percent of</u> <u>Revenue</u>	<u>Dollar Change</u>	<u>Percentage</u> <u>Change</u>
Sales and marketing	\$ 7,116	11.2 %	\$ 6,780	10.9 %	\$ 336	5.0 %
General and administrative	4,658	7.3 %	4,778	7.7 %	(120)	(2.5 %)
Research and development	1,857	2.9 %	1,791	2.9 %	66	3.7 %
Total operating expenses	<u>\$ 13,631</u>	21.4 %	<u>\$ 13,349</u>	21.5 %	<u>\$ 282</u>	2.1 %

The increase in sales and marketing expense resulted primarily from higher personnel-related costs, partly offset by the favorable impact of changes in foreign currency exchange rates. General and administrative expense and research and development expense for the nine months ended September 30, 2012 were generally consistent with the same period of the prior year.

Livestock and Poultry Diagnostics. The following table presents LPD operating expenses by functional area:

<u>Operating Expenses</u> <i>(dollars in thousands)</i>	<u>For the Nine</u> <u>Months Ended</u> <u>September 30,</u> <u>2012</u>	<u>Percent of</u> <u>Revenue</u>	<u>For the Nine</u> <u>Months Ended</u> <u>September 30,</u> <u>2011</u>	<u>Percent of</u> <u>Revenue</u>	<u>Dollar Change</u>	<u>Percentage</u> <u>Change</u>
Sales and marketing	\$ 11,416	17.8 %	\$ 11,789	16.8 %	\$ (373)	(3.2 %)
General and administrative	8,861	13.8 %	9,031	12.9 %	(170)	(1.9 %)
Research and development	8,626	13.4 %	8,754	12.5 %	(128)	(1.5 %)
Total operating expenses	<u>\$ 28,903</u>	45.1 %	<u>\$ 29,574</u>	42.3 %	<u>\$ (671)</u>	(2.3 %)

The decrease in sales and marketing expense resulted primarily from the favorable impact of changes in foreign currency exchange rates. The decrease in general and administrative expense resulted primarily from a decrease in intangible asset amortization and the favorable impact of changes in foreign currency exchange rates. The decrease in research and development expense was due primarily to lower personnel-related costs and the favorable impact of changes in foreign currency exchange rates.

Other. Operating expenses for Other increased \$1.1 million to \$13.8 million for the nine months ended September 30, 2012 due primarily to higher personnel-related costs in our OPTI Medical line of business.

Unallocated Amounts. Operating expenses that are not allocated to our operating segments decreased \$2.4 million to \$6.1 million for the nine months ended September 30, 2012 due primarily to lower personnel-related costs, lower legal and other professional fees incurred in connection with the U.S. FTC investigation and the absence of legal and other fees incurred in connection with the U.K. Office of Fair Trading (“OFT”) investigation, which concluded in November 2011. These favorable factors were partly offset by the absence of a payment related to the sale of certain raw material inventory in connection with the restructuring of our pharmaceutical business in 2008 that was recognized during the nine months ended September 30, 2011 and certain foreign exchange losses during the nine months ended September 30, 2012 compared to gains during the same period of the prior year. We estimate certain personnel-related costs and allocate the estimated expense to the operating segments. This allocation differs from the actual expense and consequently yields a difference that is reported under the caption “Unallocated Amounts.” The FTC investigation is discussed in more detail under the heading “Part II. Item 1A. Risk Factors” in this Quarterly Report on Form 10-Q.

Interest Income and Interest Expense

Interest income was \$1.4 million for the nine months ended September 30, 2012 compared to \$1.2 million for the same period of the prior year.

Interest expense was \$3.0 million for the nine months ended September 30, 2012 compared to \$2.4 million for the same period of the prior year. The increase in interest expense was due primarily to higher average balances outstanding on our Credit Facility, partly offset by lower effective interest rates.

Provision for Income Taxes

Our effective income tax rate was 31.7% for the nine months ended September 30, 2012, compared to 31.1% for the nine months ended September 30, 2011. The increase in our effective income tax rate was due primarily to federal research and development tax incentives that were not available during the nine months ended September 30, 2012 but were available during the same period of the prior year.

■ Recent Accounting Pronouncements

A discussion of recent accounting pronouncements is included in Note 2 to the condensed consolidated financial statements included in this Quarterly Report on Form 10-Q.

■ Liquidity and Capital Resources

Liquidity

We fund the capital needs of our business through cash on hand, funds generated from operations, and amounts available under our Credit Facility. At September 30, 2012 and December 31, 2011, we had \$221.4 million and \$183.9 million, respectively, of cash and cash equivalents, and working capital of \$154.9 million and \$87.3 million, respectively. Additionally, at September 30, 2012, we had remaining borrowing availability of \$67 million under our \$300 million Credit Facility. We believe that, if necessary, we could obtain additional borrowings at prevailing market interest rates to fund our growth objectives. We further believe that current cash and cash equivalents, funds generated from operations, and available borrowings will be sufficient to fund our operations, capital purchase requirements, and strategic growth needs for the next twelve months, and that these resources will be sufficient in the long term to fund our business as currently conducted.

We consider the majority of the operating earnings of certain non-U.S. subsidiaries to be indefinitely invested U.S. Changes to this position could have adverse tax consequences. We manage our worldwide cash requirements available funds among all of our subsidiaries. Our foreign cash balances are generally available without restrictions to ordinary business operations outside the U.S. Of our total cash and cash equivalents at September 30, 2012, approximately million was held by our foreign subsidiaries and was subject to material repatriation tax effects. The amount of cash and cash equivalents held by foreign subsidiaries subject to other restrictions on the free flow of funds (primarily securing various obligations) was approximately \$1.6 million. As of September 30, 2012, 38% of the cash and cash equivalents held by our foreign subsidiaries was invested in money market funds restricted to U.S. government and agency securities, 38% was held as bank deposits, 21% was invested in money market funds having investments in highly liquid investment-grade fixed-income securities, and 3% was invested in money market funds restricted to non-U.S. government securities corresponding to the investment currency. As of September 30, 2012, approximately 64% of the cash and cash equivalents held by our foreign subsidiaries was held in U.S. dollars.

Should we require more capital in the U.S. than is generated by our operations domestically, for example to fund significant discretionary activities, we could elect to repatriate future earnings from foreign jurisdictions or raise capital in the U.S. through debt or equity issuances. These alternatives could result in higher effective tax rates, increased interest expense, or other dilution of our earnings. We have borrowed funds domestically and continue to have the ability to borrow funds domestically at reasonable interest rates.

The following table presents additional key information concerning working capital:

	For the Three Months Ended				
	September 30, 2012	June 30, 2012	March 31, 2012	December 31, 2011	September 30, 2011
Days sales outstanding ⁽¹⁾	41.7	41.9	42.7	41.0	43.1
Inventory turns ⁽²⁾	1.7	1.8	1.8	1.8	1.7

⁽¹⁾ Days sales outstanding represents the average of the accounts receivable balances at the beginning and end of each quarter divided by revenue for that quarter, the result of which is then multiplied by 91.25 days.

⁽²⁾ Inventory turns represents inventory-related cost of product revenue for the 12 months preceding each quarter-end divided by the inventory balance at the end of the quarter.

Sources and Uses of Cash

The following table presents cash provided (used):

<i>(dollars in thousands)</i>	For the Nine Months Ended September 30,		
	2012	2011	Dollar Change
Net cash provided by operating activities	\$ 153,488	\$ 163,518	\$ (10,030)
Net cash used by investing activities	(41,085)	(39,304)	(1,781)
Net cash used by financing activities	(75,496)	(101,667)	26,171
Net effect of changes in exchange rates on cash	639	2,037	(1,398)
Net increase in cash and cash equivalents	\$ 37,546	\$ 24,584	\$ 12,962

Operating Activities. Cash provided by operating activities was \$153.5 million for the nine months ended September 30, 2012 as compared to \$163.5 million for the same period of the prior year. The total of net income and net non-cash charges, excluding the impact of reclassifying the tax benefit from share based compensation arrangements to a financing activity, was \$184.4 million for the nine months ended September 30, 2012 compared to \$172.4 million for the same period in 2011, resulting in incremental operating cash flows of \$12.0 million driven primarily by the increase in net income. The total of changes in operating assets and liabilities and the tax benefit from share-based compensation arrangements decreased cash by \$31.0 million and \$8.9 million for the nine months ended September 30, 2012 and 2011, respectively, resulting in an incremental decrease in cash of \$22.1 million.

The following table presents cash flows from changes in operating assets and liabilities and the tax benefit from share-based compensation arrangements:

<i>(dollars in thousands)</i>	For the Nine Months Ended September 30,		
	2012	2011	Dollar Change
Accounts receivable	\$ 255	\$ (16,181)	\$ 16,436
Inventories	(18,078)	(9,732)	(8,346)
Other assets	539	3,056	(2,517)
Accounts payable	(4,893)	8,305	(13,198)
Accrued liabilities	(3,751)	16,664	(20,415)
Deferred revenue	5,151	3,018	2,133
Tax benefit from share-based compensation arrangements	(10,182)	(14,009)	3,827
Total change in cash due to changes in operating assets and liabilities and the tax benefit from share-based compensation arrangements	\$ (30,959)	\$ (8,879)	\$ (22,080)

The increase in inventories during the nine months ended September 30, 2012 was more than the increase during the same period of the prior year due primarily to the timing of inventory receipts from third party suppliers, most significantly our Procyte[®] Dx instruments and the slides used in our chemistry instruments. Incremental cash used by accounts payable and accrued liabilities was due primarily to the timing of vendor and income tax payments during the nine months ended September 30, 2012 compared to the same period of the prior year. The amount and timing of revenues and customer payments was the primary driver of the cash used by accounts receivable during the nine months ended September 30, 2011.

We historically have experienced proportionally lower net cash flows from operating activities during the first quarter and proportionally higher cash flows from operating activities for the remainder of the year and for the annual period driven primarily by the payment of annual bonuses in connection with employee incentive programs in the first quarter following the year for which the bonuses were earned and the seasonality of vector-borne disease testing, which has historically resulted in significant increases in accounts receivable balances during the first quarter of the year.

Investing Activities. Cash used by investing activities was \$41.1 million for the nine months ended September 30, 2012 compared to \$39.3 million for the same period of the prior year. The increase in cash used by investing activities was due primarily to incremental investments during the nine months ended September 30, 2012 to expand our headquarters facility in Westbrook, Maine and our suite of business applications and related hardware. These increases were partly offset by the substantial completion in 2011 of investments in our Memphis, Tennessee facility.

Our total capital expenditure plan for 2012 is estimated to be approximately \$60 million, which includes approximately \$13 million for the expansion of our headquarters facility in Westbrook, Maine, approximately \$16 million for investments in our suite of business software applications and related hardware and approximately \$13 million for investments in our reference laboratory equipment and facilities.

Financing Activities. Cash used by financing activities was \$75.5 million for the nine months ended September 30, 2012 compared to cash used of \$101.7 million for the same period in 2011. The decrease in cash used by financing activities was due primarily to a decrease in cash used to repurchase common stock, partly offset by net payments under the Credit Facility for the nine months ended September 30, 2012 compared to net borrowings for the same period of the prior year and a decrease in cash received from the exercise of stock options and employee stock purchase plans and related tax benefits.

Net borrowing and repayment activity under our Credit Facility resulted in incremental cash used of \$35.9 million during the nine months ended September 30, 2012 compared to the same period of the prior year. At September 30, 2012, we had \$232.0 million outstanding under the Credit Facility. The general availability of funds under the Credit Facility was further reduced by \$1.0 million for a letter of credit issued related to our worker's compensation policy covering claims for the years ending 2009 through 2012. The obligations under the Credit Facility may be accelerated upon the occurrence of an event of default under the Credit Facility, which includes customary events of default including, without limitation, payment defaults, defaults in the performance of affirmative and negative covenants, the inaccuracy of representations or warranties, bankruptcy and insolvency related defaults, defaults relating to judgments, certain events related to employee pension benefit plans under the Employee Retirement Income Security Act of 1974, the failure to pay specified indebtedness, and a change of control default. The Credit Facility contains affirmative, negative and financial covenants customary for financings of this type. The negative covenants include restrictions on liens, indebtedness of subsidiaries of the Company, fundamental changes, investments, transactions with affiliates and certain restrictive agreements. The financial covenant is a consolidated leverage ratio test that requires our ratio of debt to earnings before interest, taxes, depreciation and amortization, defined as the consolidated leverage ratio under the terms of the Credit Facility, not to exceed 3-to-1. At September 30, 2012, we were in compliance with the covenants of the Credit Facility.

Cash used to repurchase shares of our common stock in the open market decreased by \$74.9 million during the nine months ended September 30, 2012 compared to the same period of the prior year. From the inception of our share repurchase program in August 1999 to September 30, 2012, we have repurchased 44.7 million shares. During the nine months ended September 30, 2012, we purchased 1.0 million shares for an aggregate cost of \$91.2 million compared to purchases of 2.2 million shares for an aggregate cost of \$166.0 million during the nine months ended September 30, 2011. We believe that the repurchase of our common stock is a favorable investment and we also repurchase to offset the dilutive effect of our share-based compensation programs. Repurchases of our common stock may vary depending upon the level of other investing activities and the share price. See Note 9 to the condensed consolidated financial statements included in this Quarterly Report on Form 10-Q for additional information about our share repurchases.

Cash proceeds from the exercise of stock options and employee stock purchase plans and the related tax benefits decreased by \$12.8 million during the nine months ended September 30, 2012 compared to the same period in 2011 due primarily to a decrease in the number of stock options exercised.

Other Commitments, Contingencies and Guarantees

Significant commitments, contingencies and guarantees at September 30, 2012 are consistent with those discussed in the section under the heading "Part 2, Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations—Liquidity and Capital Resources," and in Note 14 to the consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2011, except for the FTC investigation as described in Note 13 to the condensed consolidated financial statements included in this Quarterly Report on Form 10-Q.

Item 3. Quantitative and Qualitative Disclosures About Market Risk

For quantitative and qualitative disclosures about market risk affecting IDEXX, see the section under the heading "Part II, Item 7A. Quantitative and Qualitative Disclosures About Market Risk" of our Annual Report on Form 10-K for the year ended December 31, 2011. As of the date of this report, there have been no material changes to the market risks described in our Annual Report on Form 10-K for the year ended December 31, 2011.

Item 4. Controls and Procedures

Disclosure Controls and Procedures

Our management is responsible for establishing and maintaining disclosure controls and procedures, as defined by the Securities and Exchange Commission (the “SEC”) in its Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934 as amended (the “Exchange Act”). The term “disclosure controls and procedures,” as defined in Rules 13a-15(e) and 15d-15(e) under the Exchange Act, means controls and other procedures of an issuer that are designed to ensure that information required to be disclosed by the issuer in the reports that it files or submits under the Exchange Act is recorded, processed, summarized and reported, within the time periods specified in the SEC's rules and forms. Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information required to be disclosed by a company in the reports that it files or submits under the Exchange Act is accumulated and communicated to the company's management, including its principal executive and principal financial officers, as appropriate to allow timely decisions regarding required disclosure. 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 necessarily applies its judgment in evaluating the cost-benefit relationship of possible controls and procedures. Based on the evaluation of our disclosure controls and procedures at September 30, 2012, our chief executive officer and chief financial officer have concluded that, as of such date, our disclosure controls and procedures were effective at the reasonable assurance level.

Changes in Internal Control Over Financial Reporting

There were no changes in our internal control over financial reporting (as defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act) during the three months ended September 30, 2012 that materially affected, or are reasonably likely to materially affect, the Company's internal control over financial reporting.

PART II — OTHER INFORMATION

Item 1A. Risk Factors

Our future operating results involve a number of risks and uncertainties. Actual events or results may differ materially from those discussed in this report. Factors that could cause or contribute to such differences include, but are not limited to, the factors discussed below, as well as those discussed elsewhere in this report.

The following discussion includes five revised risk factors (“Our Biologic Products Are Complex and Difficult to Manufacture, Which Could Negatively Affect Our Ability to Supply the Market,” “Strengthening of the Rate of Exchange for the U.S. Dollar Has a Negative Effect on Our Business,” “The Resolution of the FTC Investigation into Our Marketing and Sales Practices for Companion Animal Veterinary Products and Services is Uncertain,” “Increase in Corporate Hospital Ownership and Prevalence of Buying Consortiums Could Negatively Affect Our Business” and “Risks Associated with Doing Business Internationally Could Negatively Affect Our Operating Results”) that reflect developments subsequent to the discussion of those risk factors included in our Annual Report on Form 10-K for the year ended December 31, 2011. Further, we have added one risk factor (“The Impact of a Change in the Status of One of Our Distributors on Our Results of Operations is Uncertain”) that was not included in our Annual Report on Form 10-K for the year ended December 31, 2011.

Our Failure to Successfully Execute Certain Strategies Could Have a Negative Impact on Our Growth and Profitability

The companion animal health care industry is highly competitive and we anticipate increasing levels of competition from both existing competitors and new market entrants. Our ability to maintain or enhance our growth rates and our profitability depends on our successful execution of many elements of our strategy, which include:

- Developing, manufacturing and marketing innovative new or improved in-clinic laboratory analyzers that drive sales of IDEXX VetLab[®] instruments, grow our installed base of instruments, and increase demand for related consumable products, services and accessories;
- Developing and introducing new proprietary diagnostic tests and services that provide valuable medical information to our customers and effectively differentiate our products and services from those of our competitors;
- Increasing the value to our customers of our companion animal products and services by enhancing the integration of these products and the management of diagnostic information derived from our products;

- Providing our veterinary customers with the medical and business tools, information and resources that enable them to grow their practices through increased pet visits and enhanced practice of real-time care;
- Achieving cost improvements in our worldwide network of laboratories by implementing global best practices including lean processing techniques, incorporating technological enhancements including laboratory automation and a global laboratory information management system, employing purchasing strategies to maximize leverage of our global scale, increasing the leverage of existing infrastructure and consolidating testing in high volume laboratory hubs;
- Achieving cost improvements in the manufacture and service of our in-clinic laboratory analyzers by employing the benefits of economies of scale in both negotiating supply contracts and leveraging manufacturing overhead, and by improving reliability of our instruments;
- Expanding our served market and growing our market share by strengthening our sales and marketing activities both within the U.S. and in geographies outside of the U.S.;
- Identifying, completing and integrating acquisitions that enhance our existing businesses or create new business or geographic areas for us; and
- Developing and implementing new technology and licensing strategies.

If we are unsuccessful in implementing and executing on some or all of these strategies, our rate of growth or profitability may be negatively impacted.

Our Dependence on a Limited Number of Suppliers Could Limit Our Ability to Sell Certain Products or Reduce Our Profitability

We currently purchase many products and materials from sole or single sources. Some of the products that we purchase from these sources are proprietary and, therefore, cannot be readily or easily replaced by alternative sources. These products include our Catalyst Dx[®] and VetTest[®] consumables; ProCyte Dx[®] hematology, IDEXX VetAutoread[™] hematology, VetLyte[®] electrolyte, IDEXX VetLab[®] UA[™] urinalysis, VetTest[®] chemistry, and Coag Dx[™] blood coagulation analyzers and related consumables and accessories; image capture plates used in our digital radiography systems; and certain components and raw materials used in our SNAP[®] rapid assay devices, water testing products, livestock and poultry diagnostic tests, dairy testing products and LaserCyte[®] hematology analyzers. To mitigate risks associated with sole and single source suppliers, we seek when possible to enter into long-term contracts that ensure an uninterrupted supply of products at predictable prices. However, some suppliers decline to enter into long-term contracts and we are required to purchase products on a purchase order basis. There can be no assurance that suppliers with which we do not have contracts will continue to supply our requirements for products, that suppliers with which we do have contracts will always fulfill their obligations under these contracts, or that any of our suppliers will not experience disruptions in their ability to supply our requirements for products. In cases where we purchase sole and single source products or components under purchase orders, we are more susceptible to unanticipated cost increases or changes in other terms of supply. In addition, under some contracts with suppliers we have minimum purchase obligations and our failure to satisfy those obligations may result in loss of some or all of our rights under these contracts or require us to compensate the supplier. If we are unable to obtain adequate quantities of sole and single source products in the future, we may be unable to supply the market, which could have a material adverse effect on our results of operations.

Our Biologic Products Are Complex and Difficult to Manufacture, Which Could Negatively Affect Our Ability to Supply the Market

Many of our rapid assay, livestock and poultry diagnostic, water and dairy products are biologic products, which are products that include materials from living organisms, such as antibodies, cells and sera. Manufacturing biologic products is highly complex due to the inherent variability of biological input materials and to the difficulty of controlling the interactions of these materials with other components of the products, with samples and with the environment. There can be no assurance that we will be able to maintain adequate sources of biological materials or that we will be able to consistently manufacture biologic products that satisfy applicable product release criteria. Further, products that meet release criteria at the time of manufacture may fall out of specification while in customer inventory, which could necessitate field actions that would require us to incur expenses associated with recalling products and providing customers with new products, and could damage customer relations. Our inability to produce or obtain necessary biological materials or to successfully manufacture biologic products that incorporate such materials could result in our inability to supply the market with these products, which could have a material adverse effect on our results of operations.

A Weak Economy Could Result in Reduced Demand for Our Products and Services or Increased Customer Credit Risk

A substantial percentage of our sales are made worldwide to the companion animal veterinary market. Demand for our companion animal diagnostic products and services is driven in part by the number of patient visits with their respective owners to veterinary hospitals and the practices of veterinarians with respect to the recommendations for diagnostic testing, as well as the pet owner compliance with these recommendations. Economic weakness in our significant markets in recent years has caused and could continue to cause pet owners to skip or defer visits to veterinary hospitals or could affect their willingness to approve certain diagnostic tests, comply with a treatment plan or, even more fundamentally, continue to own a pet. In addition, concerns about the financial resources of pet owners could cause veterinarians to be less likely to recommend certain diagnostic tests and concerns about the economy may cause veterinarians to defer purchasing capital items such as our instruments and systems. A decline in patient visits to the hospital, in the willingness of pet owners to treat certain health conditions or approve certain tests, in pet ownership in general, or in the inclination of veterinarians to recommend certain tests or make capital purchases could result in a decrease in sales of diagnostic products and services, which could have a material adverse effect on our results of operations.

Demand for our water products is driven in part by the availability of funds at the government laboratories, water utilities and private certified laboratories that utilize our products. Availability of funds also affects demand by the government laboratories and cattle, swine and poultry producers that utilize our livestock and poultry diagnostic products, and by users of our human point-of-care diagnostic instruments. Economic weakness in our markets has caused and could continue to cause our customers to reduce their investment in such testing, which could have a material adverse effect on our results of operations.

In all of our markets, a weak economy may also cause deterioration in the financial condition of our distributors and customers, which could inhibit their ability to pay us amounts owed for products delivered or services provided in a timely fashion or at all.

Strengthening of the Rate of Exchange for the U.S. Dollar Has a Negative Effect on Our Business

Any strengthening of the rate of exchange for the U.S. dollar against non-U.S. currencies, and in particular the Euro, British pound, Canadian dollar, Japanese yen and Australian dollar, adversely affects our results, as it reduces the dollar value of sales that are made in those currencies and reduces the profits on products manufactured or sourced in U.S. dollars and exported to international markets. For both the three and nine months ended September 30, 2012, approximately 25% of our revenue was derived from products manufactured in the U.S. and sold internationally in local currencies compared to 25% and 26%, respectively, for the three and nine months ended September 30, 2011. To mitigate such foreign currency exposure, we utilize non-speculative forward currency exchange contracts. A strengthening U.S. dollar could also negatively impact the ability of customers outside the U.S. to pay for purchases denominated in U.S. dollars.

Various Government Regulations Could Limit or Delay Our Ability to Market and Sell Our Products

In the U.S., the manufacture and sale of our products are regulated by agencies such as the United States Department of Agriculture (“USDA”), the U.S. Food and Drug Administration (“FDA”) and the U.S. Environmental Protection Agency (“EPA”). Our infectious disease diagnostic tests for animal health applications, including most rapid assay canine and feline SNAP[®] tests and livestock and poultry diagnostic tests, must be approved by the USDA prior to sale in the U.S. Our water testing products must be approved by the EPA before they can be used by customers in the U.S. as a part of a water quality monitoring program required by the EPA. Our dairy testing products require approval by the FDA prior to sale in the U.S. The manufacture and sale of our OPTI[®] line of human point-of-care electrolytes and blood gas analyzers are regulated by the FDA and these products require approval by the FDA before they may be sold commercially in the U.S. The manufacture and sale of our products are subject to similar and sometimes more stringent laws in many foreign countries. Any failure to comply with legal and regulatory requirements relating to the manufacture and sale of our products in the U.S. or in other countries could result in fines and sanctions against us or suspensions or discontinuations of our ability to manufacture or sell our products, which could have a material adverse effect on our results of operations. In addition, delays in obtaining regulatory approvals for new products or product upgrades could have a negative impact on our growth and profitability.

The Resolution of the FTC Investigation into Our Marketing and Sales Practices for Companion Animal Veterinary Products and Services is Uncertain

As discussed in our Annual Report on Form 10-K for the year ended December 31, 2011, in January 2010, we received a letter from the U.S. Federal Trade Commission (“FTC”), stating that it was conducting an investigation to determine whether IDEXX or others have engaged in, or are engaging in, unfair methods of competition in violation of Section 5 of the Federal Trade Commission Act (“FTC Act”), through pricing or marketing policies for companion animal veterinary products and services, including but not limited to exclusive dealing or tying arrangements with distributors or end-users of those products or services. The letter requested that we preserve all materials potentially relevant to this investigation. The letter stated that the FTC had not concluded that IDEXX or anyone else had violated Section 5 of the FTC Act. We received subpoenas from the FTC on April 15, 2010 and August 8, 2011, requesting that we provide the FTC with documents and information relevant to this investigation and we have cooperated fully with the FTC in its investigation.

Based on discussions with the FTC staff, we determined that their primary concern was that our relationships with the three largest U.S. distributors of companion animal products, referred to as the national distributors, limited access by our competitors to companion animal veterinary practices. We also determined that their concerns could be addressed if one of these three national distributors was free to carry competitive products. In response to these concerns, in August 2012 we announced our intention to seek to modify our agreement with one of these distributors to eliminate our competitive products policy, which permits us to discontinue sale of our products in a particular product category to the distributor if the distributor sells competitive products in that product category. On September 28, 2012, we entered into such a modified agreement with MWI Veterinary Supply, Inc. (“MWI”) that will become effective January 1, 2013. Under this modified agreement, MWI will be permitted to carry any competitive products without restriction or potential negative consequence. This agreement was reviewed by the FTC staff prior to its execution by IDEXX and MWI to ensure that the agreement met the staff’s objective of ensuring that one national distributor had the right to carry competitive products without limitation.

We are currently negotiating the terms of a consent order with the FTC staff that we expect will require, among other things, that at least one national distributor will not be exclusive to IDEXX for the duration of the consent order, which is expected to be ten years. If successfully negotiated with the staff, the consent order would need to be approved by the FTC Commissioners prior to effectiveness. We cannot provide any assurance that we will be able to successfully negotiate the consent order with the staff, or that the FTC Commissioners will approve the consent order.

If a consent order is not approved by the FTC Commissioners, we cannot provide any assurances that the FTC will not determine to file a complaint against IDEXX in the administrative law court within the FTC at some point in the future, or that we would be successful defending ourselves in such a proceeding. Were we to be unsuccessful in defending this proceeding and any applicable appeals, we could be subject to restrictions on certain of our marketing and sales practices, including on the terms included in our agreements with certain of our U.S. distributors. While we cannot be certain about what prospective remedies would be sought by the FTC in any such proceeding, we believe that any required changes in our marketing or sales practices would not have a material adverse effect on our financial statements.

We continue to believe that our marketing and sales practices for companion animal veterinary products and services do not violate applicable antitrust laws. We have chosen to enter into the modified agreement with MWI and negotiate a consent order with the FTC staff because we believe this course will help us avoid long and costly litigation, and that our business will not be materially adversely affected.

The Impact of a Change in the Status of One of Our Distributors on Our Results of Operations is Uncertain

The modification of our agreement with MWI to eliminate our competitive products policy, as described in the risk factor above, will result in one or more of our competitors selling products through MWI, which we expect will increase the field sales resources used by those competitors to sell their products. Under the modified agreement with MWI, we will provide lower compensation to MWI on sales of our products since we will no longer receive the benefits of MWI’s exclusive focus on our products. We expect to reinvest savings from this lower rate of compensation in other sales and marketing resources and the selling efforts of our other distributors. We believe that the reallocation of these sales resources will help mitigate the potential effects of the loss of exclusive focus of MWI and the additional field sales resources used by our competitors. However, there can be no assurances that we will be able to fully mitigate the competitive effects of the changes in the nature of our agreement with MWI. Any reduction in the relative effectiveness of our overall selling efforts could have an adverse effect on our results of operations, which we do not believe would be material.

Our Success Is Heavily Dependent Upon Our Proprietary Technologies

We rely on a combination of patent, trade secret, trademark and copyright laws to protect our proprietary rights. If we do not have adequate protection of our proprietary rights, our business may be affected by competitors who utilize substantially equivalent technologies that compete with us.

We cannot ensure that we will obtain issued patents, that any patents issued or licensed to us will remain valid, or that any patents owned or licensed by us will provide protection against competitors with similar technologies. Even if our patents cover products sold by our competitors, the time and expense of litigating to enforce our patent rights could be substantial, and could have a material adverse effect on our results of operations. In addition, expiration of patent rights could result in substantial new competition in the markets for products previously covered by those patent rights.

In the past, we have received notices claiming that our products infringe third-party patents and we may receive such notices in the future. Patent litigation is complex and expensive, and the outcome of patent litigation can be difficult to predict. We cannot ensure that we will win a patent litigation case or negotiate an acceptable resolution of such a case. If we lose, we may be stopped from selling certain products and/or we may be required to pay damages and/or ongoing royalties as a result of the lawsuit. Any such adverse result could have a material adverse effect on our results of operations.

Distributor Purchasing Patterns Could Negatively Affect Our Operating Results

We sell many of our products, including substantially all of the rapid assays and instrument consumables sold in the U.S., through distributors. Distributor purchasing patterns can be unpredictable and may be influenced by factors unrelated to the end-user demand for our products. In addition, our agreements with U.S. distributors may generally be terminated by the distributors for any reason on 60 days notice. Because significant product sales are made to a limited number of distributors, the unanticipated loss of a distributor or unanticipated changes in the frequency, timing or size of distributor purchases, could have a negative effect on our results of operations.

Distributors of veterinary products have entered into business combinations resulting in fewer distribution companies. Consolidation within distribution channels increases our customer concentration level, which could increase the risks described in the preceding paragraph. See “Part I. Item 1 Business – Marketing and Distribution” in our Annual Report on Form 10-K for the year ended December 31, 2011 for additional information.

Increased Competition and Technological Advances by Our Competitors Could Negatively Affect Our Operating Results

We face intense competition within the markets in which we sell our products and services and we expect that future competition may become even more intense. Competition could negatively affect our sales and profitability in a number of ways. New competitors may enter our markets and new or existing competitors may introduce new and competitive products and services, which could be superior to our products and services. Some of our competitors and potential competitors may choose to differentiate themselves by offering similar products and services to ours at lower sales prices, which could have a material adverse effect on our results of operations through loss of market share or a decision to lower our own sales prices to remain competitive. In addition, multiple competitors could bundle product and service offerings through co-marketing or other arrangements, which could enhance their ability to compete with our broad product and service offering. Some of our competitors and potential competitors, including large diagnostic and pharmaceutical companies, have substantially greater financial resources than us, and greater experience in manufacturing, marketing, research and development and obtaining regulatory approvals than we do.

Changes in Testing Patterns Could Negatively Affect Our Operating Results

The market for our companion animal and livestock and poultry diagnostic tests and our dairy and water testing products could be negatively impacted by a number of factors impacting testing practices. The introduction or broad market acceptance of vaccines or preventatives for the diseases and conditions for which we sell diagnostic tests and services could result in a decline in testing. Changes in accepted medical protocols regarding the diagnosis of certain diseases and conditions could have a similar effect. Eradication or substantial declines in the prevalence of certain diseases also could lead to a decline in diagnostic testing for such diseases. Our livestock and poultry products business in particular is subject to fluctuations resulting from changes in disease prevalence. In addition, changes in government regulations or in the availability of government funds available for monitoring programs could negatively affect sales of our products that are driven by compliance testing, such as our livestock and poultry, dairy and water products. Declines in testing for any of the reasons described, along with lost opportunities associated with a reduction in veterinary visits, could have a material adverse effect on our results of operations.

Effective January 1, 2009, the age at which healthy cattle to be slaughtered are required to be tested for bovine spongiform encephalopathy (“BSE” or “mad cow disease”) in the European Union was increased from 30 months to 48 months, which reduced the population of cattle tested by approximately 30%. In February 2011, the European Union’s Standing Committee on the Food Chain and Animal Health agreed to allow its member states to further raise the recommended testing age to 72 months, effective July 1, 2011, which is further reducing the population of cattle tested. The demand for our BSE testing products has been negatively impacted as a result of these regulatory changes, and could be further impacted by further changes that could be made in the future.

Increase in Corporate Hospital Ownership and Prevalence of Buying Consortiums Could Negatively Affect Our Business

An increasing percentage of veterinary hospitals in the U.S. is owned by corporations that are in the business of acquiring veterinary hospitals and/or opening new veterinary hospitals nationally or regionally. Major corporate hospital owners in the U.S. include VCA Antech, Inc., National Veterinary Associates and Banfield Pet Hospital, each of which is currently a customer of IDEXX. A similar trend exists in the U.K. and may in the future also develop in other countries. Furthermore, an increasing percentage of individually owned veterinary hospitals in the U.S. are participating in buying consortiums. Corporate owners of veterinary hospitals and buying consortiums often seek to improve profitability by leveraging the buying power they derive from their scale to obtain favorable pricing from suppliers, which could have a negative impact on our results of operations. While we have strong supplier relationships with several corporate hospital groups and buying consortiums that we believe are positive for our business, decisions by larger corporate owners and buying consortiums, in particular Banfield Pet Hospital, to shift their purchasing of products and services away from us and to a competitor would have a negative impact on our results of operations, which could be material. In addition, certain corporate owners, most notably VCA Antech, our primary competitor in the U.S. and Canadian markets for veterinary reference laboratory diagnostic services, also operate reference laboratories that serve both their hospitals and unaffiliated hospitals. Any hospitals acquired by these companies generally use their reference laboratory services almost exclusively and shift a large portion of their testing from in-clinic testing to their reference laboratories. Furthermore, because these companies compete with us in the reference laboratory services marketplace, hospitals acquired by these companies may cease to be customers or potential customers of our other companion animal products and services, which would cause our sales of these products and services to decline.

Our Limited Experience and Small Scale in the Human Point-of-Care Market Could Inhibit Our Success in this Market

We have limited experience in the human point-of-care medical diagnostics market and we operate at a small scale in this market. This market differs in many respects from the veterinary diagnostic market. Significant differences include the impact of third party reimbursement on diagnostic testing, more extensive regulation, greater product liability risks, larger competitors, a more segmented customer base and more rapid technological innovation. Our limited experience and small scale in the human point-of-care medical diagnostics market could negatively affect our ability to successfully manage the risks and features of this market that differ from the veterinary diagnostic market. There can be no assurance that we will be successful in achieving growth and profitability in the human point-of-care medical diagnostics market comparable to the results we have achieved in the veterinary diagnostic market.

Risks Associated with Doing Business Internationally Could Negatively Affect Our Operating Results

For both the three months and nine months ended September 30, 2012, approximately 41% of our revenue was attributable to sales of products and services to customers outside the U.S., compared to 42% for both the three and nine months ended September 30, 2011. Various possible risks associated with foreign operations may impact our international sales, including disruptions in transportation of our products, the differing product and service needs of foreign customers, difficulties in building and managing foreign operations, import/export duties and licensing requirements, natural disasters and unexpected regulatory and economic or political changes in foreign markets. Further, prices that we charge to foreign customers may be different than the prices we charge for the same products in the U.S. due to competitive, market or other factors. Our results of operations are also susceptible to changes in foreign currency exchange rates. As a result, the mix of domestic and international sales in a particular period could have a material impact on our results of operations for that period.

Our Operations are Vulnerable to Interruption as a Result of Natural and Man-Made Disasters or System Failures

The operation of all of our facilities may be vulnerable to interruption as a result of natural and man-made disasters, interruptions in power supply or other system failures. While we maintain plans to continue business under such circumstances, there can be no assurance that such plans will be successful in fully or partially mitigating the effects of such events.

We manufacture many of our significant companion animal products, including our rapid assay devices and certain instruments, many of our water testing products and certain of our livestock and poultry testing products, at a single facility in Westbrook, Maine. Certain of our companion animal products, as well as our human point-of-care products, are manufactured in Roswell, Georgia. We also manufacture certain of our livestock and poultry testing products in Bern, Switzerland and Montpellier, France. In addition, we maintain major distribution facilities in North America and in the Netherlands and major reference laboratories in Memphis, Tennessee; Ludwigsburg, Germany; Sacramento, California; Elmhurst, Illinois; North Grafton, Massachusetts; East Brisbane, Australia; Markham, Ontario; and Wetherby, the United Kingdom. Therefore, interruption of operations at any of these facilities could have a material adverse effect on our results of operations.

We rely on several information systems throughout our company to keep financial records, process customer orders, manage inventory, process shipments to customers and operate other critical functions. If we were to experience a system disruption that impacts any of our critical functions, it could result in the loss of sales and customers, financial misstatement and significant incremental costs, which could adversely affect our business.

We maintain property and business interruption insurance to insure against the financial impact of certain events of this nature. However, this insurance may be insufficient to compensate us for the full amount of any losses that we may incur. In addition, such insurance will not compensate us for the long-term competitive effects of being out of the market for the period of any interruption in operations.

We Could Be Subject to Class Action Litigation Due to Stock Price Volatility, which, if it Occurs, Could Result in Substantial Costs or Large Judgments Against Us

The market for our common stock may experience extreme price and volume fluctuations, which may be unrelated or disproportionate to our operating performance or prospects. Securities class action litigation has often been brought against companies following periods of volatility in the market prices of their securities. We may be the target of similar litigation in the future. Securities litigation could result in substantial costs and divert our management's attention and resources, which could have a negative effect on our business, operating results and financial condition.

If Our Quarterly or Annual Results of Operations Fluctuate, This Fluctuation May Cause Our Stock Price to Decline, Resulting in Losses to You

Our prior operating results have fluctuated due to a number of factors, including seasonality of certain product lines; changes in our accounting estimates; the impact of acquisitions; timing of distributor purchases, product launches, operating expenditures, changes in foreign currency exchange rates, litigation and claim-related expenditures; changes in competitors' product offerings; changes in the economy affecting consumer spending; and other matters. Similarly, our future operating results may vary significantly from quarter to quarter or year to year due to these and other factors, many of which are beyond our control. If our operating results or projections of future operating results do not meet the expectations of market analysts or investors in future periods, our stock price may fall.

Future Operating Results Could Be Negatively Affected by the Resolution of Various Uncertain Tax Positions and by Potential Changes to Tax Incentives

In the ordinary course of our business, there are many transactions and calculations where the ultimate tax determination is uncertain. Significant judgment is required in determining our worldwide provision for income taxes. We periodically assess our exposures related to our worldwide provision for income taxes and believe that we have appropriately accrued taxes for contingencies. Any reduction of these contingent liabilities or additional assessments would increase or decrease income, respectively, in the period such determination was made. Our income tax filings are regularly under audit by tax authorities and the final determination of tax audits could be materially different than that which is reflected in historical income tax provisions and accruals. Additionally, we benefit from certain tax incentives offered by various jurisdictions. If we are unable to meet the requirements of such incentives, or if they expire or are renewed at less favorable terms, our inability to realize these benefits could have a material negative effect on future earnings.

Restrictions in Our Credit Facility or Our Inability to Obtain Financing on Favorable Terms May Limit Our Activities

Our ability to satisfy our obligations under our unsecured revolving credit facility (“Credit Facility”) depends on our future operating performance and on economic, financial, competitive and other factors beyond our control. Our business may not generate sufficient cash flows to meet these obligations or generate sufficient levels of earnings to satisfy the applicable affirmative, negative and financial covenants. Our failure to comply with these covenants and the other terms of the Credit Facility could result in an event of default and acceleration of our obligations under the Credit Facility, which may require us to seek additional financing or restructure existing debt and possibly on terms not deemed favorable.

We fund our operations, capital purchase requirements and strategic growth needs through cash on hand, funds generated from operations and amounts available under our Credit Facility. If we were unable to obtain financing on favorable terms, we could face restrictions that would limit our ability to execute certain strategies, which could have an adverse effect on our revenue growth and profitability.

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

During the three months ended September 30, 2012, we repurchased shares of common stock as described below:

Period	Total Number of Shares Purchased (a)	Average Price Paid per Share (b)	Total Number of Shares Purchased as Part of Publicly Announced Plans or Programs (c)	Maximum Number of Shares that May Yet Be Purchased Under the Plans or Programs (d)
July 1 to July 31, 2012	19,319	\$ 87.61	18,300	3,717,026
August 1 to August 31, 2012	220,800	91.62	220,800	3,496,226
September 1 to September 30, 2012	146,693	97.77	146,400	3,349,826
Total	<u>386,812</u>	\$ 93.75	<u>385,500</u>	3,349,826

Our board of directors has approved the repurchase of up to 48 million shares of our common stock in the open market or in negotiated transactions. The plan was approved and announced on August 13, 1999, and subsequently amended on October 4, 1999, November 16, 1999, July 21, 2000, October 20, 2003, October 12, 2004, October 12, 2005, February 14, 2007, February 13, 2008, February 10, 2010 and October 12, 2011 and does not have a specified expiration date. There were no other repurchase plans outstanding during the three months ended September 30, 2012, and no repurchase plans expired during the period. Repurchases of 385,500 shares were made during the three months ended September 30, 2012 in transactions made pursuant to our repurchase plan.

During the three months ended September 30, 2012, we received 1,312 shares of our common stock that were surrendered by employees in payment for the minimum required withholding taxes due on the vesting of restricted stock units and settlement of deferred stock units. In the above table, these shares are included in columns (a) and (b), but excluded from columns (c) and (d). These shares do not reduce the number of shares that may be purchased under the repurchase plan.

Item 6. Exhibits

<u>Exhibit No.</u>	<u>Description</u>
31.1	Certification of Principal Executive Officer pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
31.2	Certification of Principal Financial Officer pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
32.1	Certification of Chief Executive Officer 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 Financial Officer pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
101.INS	XBRL Instance Document.
101.SCH	XBRL Taxonomy Extension Schema Document.
101.CAL	XBRL Taxonomy Extension Calculation Linkbase Document.
101.DEF	XBRL Taxonomy Extension Definition Linkbase Document.
101.LAB	XBRL Taxonomy Extension Label Linkbase Document.
101.PRE	XBRL Taxonomy Extension Presentation Linkbase Document.

SIGNATURES

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

IDEXX LABORATORIES, INC.

Date: October 19, 2012

/s/ Merilee Raines

Merilee Raines
Executive Vice President, Chief Financial Officer
and Treasurer
(Principal Financial Officer)

Exhibit Index

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**CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002**

I, Jonathan W. Ayers, certify that:

- 1) I have reviewed this report on Form 10-Q for the quarter ended September 30, 2012 of IDEXX Laboratories, Inc.;
- 2) Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
- 3) Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
- 4) The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report, based on such evaluation; and
 - d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
- 5) The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: October 19, 2012

/s/ Jonathan W. Ayers
Jonathan W. Ayers, Chairman,
President and Chief Executive Officer
(Principal Executive Officer)

**CERTIFICATION PURSUANT TO
RULES 13a-14(a) AND 15d-14(a) UNDER THE SECURITIES EXCHANGE ACT OF 1934,
AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002**

I, Merilee Raines, certify that:

- 1) I have reviewed this report on Form 10-Q for the quarter ended September 30, 2012 of IDEXX Laboratories, Inc.;
- 2) Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
- 3) Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
- 4) The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a) Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c) Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report, based on such evaluation; and
 - d) Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
- 5) The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: October 19, 2012

/s/ Merilee Raines
Merilee Raines
Executive Vice President, Chief Financial Officer
and Treasurer
(Principal Financial Officer)

**CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED BY
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002**

In connection with the report on Form 10-Q of IDEXX Laboratories, Inc. (the “Company”) for the quarter ended September 30, 2012 as filed with the Securities and Exchange Commission on the date hereof (the “Report”), the undersigned certifies, pursuant to 18 U.S.C. Section 1350, as adopted by Section 906 of the Sarbanes-Oxley Act of 2002, that:

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

October 19, 2012

/s/ Jonathan W. Ayers

Jonathan W. Ayers, Chairman,
President and Chief Executive Officer

A signed original of this written statement required by Section 906, has been provided to IDEXX Laboratories, Inc. and will be retained by IDEXX Laboratories, Inc. and furnished to the Securities and Exchange Commission or its staff upon request.

**CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED BY
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002**

In connection with the report on Form 10-Q of IDEXX Laboratories, Inc. (the "Company") for the quarter ended September 30, 2012 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), the undersigned certifies, pursuant to 18 U.S.C. Section 1350, as adopted by Section 906 of the Sarbanes-Oxley Act of 2002, that:

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

October 19, 2012

/s/ Merilee Raines

Merilee Raines

Executive Vice President, Chief Financial Officer
and Treasurer

A signed original of this written statement required by Section 906, has been provided to IDEXX Laboratories, Inc. and will be retained by IDEXX Laboratories, Inc. and furnished to the Securities and Exchange Commission or its staff upon request.