ALLERGAN INC
Form 10-Q
May 06, 2011
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UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 10-Q

(Mark One)

x **QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**
For the quarterly period ended March 31, 2011

OR

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

Commission File Number 1-10269

Allergan, Inc.

(Exact Name of Registrant as Specified in its Charter)

Delaware	**95-1622442**
(State or Other Jurisdiction of Incorporation or Organization)	(I.R.S. Employer Identification No.)
2525 Dupont Drive	**92612**
Irvine, California	(Zip Code)
(Address of Principal Executive Offices)	

(714) 246-4500

(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 x 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 x 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 x	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 x

As of May 2, 2011, there were 307,511,888 shares of common stock outstanding (including 2,729,091 shares held in treasury).

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PART I FINANCIAL INFORMATION

Item 1. ***Financial Statements***

ALLERGAN, INC.

UNAUDITED CONDENSED CONSOLIDATED STATEMENTS OF EARNINGS

(in millions, except per share amounts)

	Three months ended	
	March 31, 2011	March 31, 2010
Revenues:		
Product net sales	$ 1,252.8	$ 1,105.8
Other revenues	18.4	48.9
Total revenues	1,271.2	1,154.7
Operating costs and expenses:		
Cost of sales (excludes amortization of acquired intangible assets)	183.3	170.2
Selling, general and administrative	589.5	473.8
Research and development	197.7	222.7
Amortization of acquired intangible assets	32.5	37.1
Intangible asset impairment	16.1	
Restructuring charges	4.6	0.6
Operating income	247.5	250.3
Non-operating income (expense):		
Interest income	2.3	1.3
Interest expense	(24.7)	(16.6)
Other, net	(9.9)	(3.0)
	(32.3)	(18.3)
Earnings before income taxes	215.2	232.0
Provision for income taxes	56.4	63.0
Net earnings	158.8	169.0
Net earnings attributable to noncontrolling interest	0.5	1.1
Net earnings attributable to Allergan, Inc.	$ 158.3	$ 167.9
Earnings per share attributable to Allergan, Inc. stockholders:		
Basic	$ 0.52	$ 0.55
Diluted	$ 0.51	$ 0.55

See accompanying notes to unaudited condensed consolidated financial statements.

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ALLERGAN, INC.

UNAUDITED CONDENSED CONSOLIDATED BALANCE SHEETS

(in millions, except share data)

	December 31, March 31, 2011	December 31, December 31, 2010
ASSETS		
Current assets:		
Cash and equivalents	$ 2,530.8	$ 1,991.2
Short-term investments	249.7	749.1
Trade receivables, net	671.6	647.3
Inventories	239.2	229.4
Other current assets	393.1	376.7
Total current assets	4,084.4	3,993.7
Investments and other assets	258.5	261.4
Deferred tax assets	242.6	217.8
Property, plant and equipment, net	788.5	800.6
Goodwill	2,045.3	2,038.6
Intangibles, net	951.4	996.0
Total assets	$ 8,370.7	$ 8,308.1
LIABILITIES AND EQUITY		
Current liabilities:		
Notes payable	$ 36.3	$ 28.1
Convertible notes	648.9	642.5
Accounts payable	200.5	222.5
Accrued compensation	118.8	182.4
Other accrued expenses	471.1	436.8
Income taxes		16.1
Total current liabilities	1,475.6	1,528.4
Long-term debt	1,529.5	1,534.2
Other liabilities	477.5	464.4
Commitments and contingencies		
Equity:		
Allergan, Inc. stockholders equity:		
Preferred stock, $.01 par value; authorized 5,000,000 shares; none issued		
Common stock, $.01 par value; authorized 500,000,000 shares; issued 307,512,000 shares as of March 31, 2011 and December 31, 2010	3.1	3.1
Additional paid-in capital	2,828.8	2,815.5
Accumulated other comprehensive loss	(131.6)	(152.9)
Retained earnings	2,342.3	2,225.9
	5,042.6	4,891.6
Less treasury stock, at cost (2,492,000 shares as of March 31, 2011 and 1,987,000 shares as of December 31, 2010)	(176.9)	(133.9)
Total stockholders equity	4,865.7	4,757.7
Noncontrolling interest	22.4	23.4

Total equity		4,888.1		4,781.1
Total liabilities and equity	$	8,370.7	$	8,308.1

See accompanying notes to unaudited condensed consolidated financial statements.

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ALLERGAN, INC.

UNAUDITED CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS

(in millions)

	Three months ended	
	March 31, 2011	March 31, 2010
Cash flows from operating activities:		
Net earnings	$ 158.8	$ 169.0
Non-cash items included in net earnings:		
Depreciation and amortization	63.4	66.3
Amortization of original issue discount and debt issuance costs	7.5	6.9
Amortization of net realized gain on interest rate swap	(0.3)	(0.3)
Deferred income tax benefit	(26.6)	(12.1)
(Gain) loss on disposal and impairment of assets	(1.5)	0.4
Unrealized loss on derivative instruments	6.9	0.7
Expense of share-based compensation plans	20.8	18.2
Intangible asset impairment	16.1	
Restructuring charges	4.6	0.6
Changes in operating assets and liabilities:		
Trade receivables	(14.2)	11.6
Inventories	(6.7)	(3.1)
Other current assets	(4.7)	(22.9)
Other non-current assets	(5.4)	(2.7)
Accounts payable	(21.6)	34.1
Accrued expenses	(53.7)	(59.3)
Income taxes	(16.1)	(21.8)
Other liabilities	10.9	(12.3)
Net cash provided by operating activities	138.2	173.3
Cash flows from investing activities:		
Purchases of short-term investments	(149.9)	
Acquisition, net of cash acquired		(63.7)
Additions to property, plant and equipment	(17.1)	(12.5)
Additions to capitalized software	(3.3)	(2.9)
Contractual purchase price adjustment to prior acquisition		(1.7)
Proceeds from maturities of short-term investments	649.3	
Proceeds from sale of property, plant and equipment	0.2	
Net cash provided by (used in) investing activities	479.2	(80.8)
Cash flows from financing activities:		
Dividends to stockholders	(15.3)	(15.1)
Payments to acquire treasury stock	(162.9)	(59.6)
Payment of contingent consideration	(2.9)	
Net borrowings (repayments) of notes payable	8.2	(3.5)
Sale of stock to employees	80.4	36.0
Excess tax benefits from share-based compensation	4.6	
Net cash used in financing activities	(87.9)	(42.2)

Effect of exchange rate changes on cash and equivalents	10.1	(7.7)
Net increase in cash and equivalents	539.6	42.6
Cash and equivalents at beginning of period	1,991.2	1,947.1
Cash and equivalents at end of period	$ 2,530.8	$ 1,989.7
Supplemental disclosure of cash flow information		
Cash paid for:		
Interest (net of amount capitalized)	$ 11.9	$ 1.2
Income taxes, net of refunds	$ 93.2	$ 104.6

See accompanying notes to unaudited condensed consolidated financial statements.

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

Note 1: Basis of Presentation

In the opinion of management, the accompanying unaudited condensed consolidated financial statements contain all adjustments necessary (consisting only of normal recurring accruals) to present fairly the financial information contained therein. These statements do not include all disclosures required by accounting principles generally accepted in the United States of America (GAAP) for annual periods and should be read in conjunction with the Company s audited consolidated financial statements and related notes for the year ended December 31, 2010. The Company prepared the unaudited condensed consolidated financial statements following the requirements of the Securities and Exchange Commission for interim reporting. As permitted under those rules, certain footnotes or other financial information that are normally required by GAAP can be condensed or omitted. The results of operations for the three month period ended March 31, 2011 are not necessarily indicative of the results to be expected for the year ending December 31, 2011 or any other period(s).

Reclassifications

Certain reclassifications of prior year amounts have been made to conform with the current year presentation.

Recently Adopted Accounting Standards

In December 2010, the Financial Accounting Standards Board (FASB) issued an accounting standards update that provides guidance on the recognition and classification of the annual fee imposed by the Patient Protection and Affordable Care Act as amended by the Health Care and Education Affordability Reconciliation Act on pharmaceutical companies that sell branded prescription drugs or biologics to specified government programs in the United States. Under this guidance, the annual fee should be estimated and recognized in full as a liability upon the first qualifying sale with a corresponding deferred cost that is amortized to operating expense using a straight-line method of allocation unless another method better allocates the fee over the calendar year in which it is payable. The annual fee ranges from $2.5 billion to $4.1 billion for all affected entities in total, a portion of which will be allocated to the Company on the basis of the amount of its branded prescription drug sales for the preceding year as a percentage of the industry s branded prescription drug sales for the same period. The annual fee is not deductible for federal income tax purposes. This guidance became effective for calendar years beginning after December 31, 2010. The Company adopted the provisions of the guidance in the first quarter of 2011 and currently estimates the annual fee for 2011 to be approximately $20.4 million.

In December 2010, the FASB issued an accounting standards update that requires an entity to perform Step 2 of the goodwill impairment test for its reporting units with a zero or a negative carrying amount if there are qualitative factors indicating that it is more likely than not that a goodwill impairment exists. This guidance became effective for fiscal years beginning after December 15, 2010 and was applied as a change in accounting principle with any impairment recorded as a cumulative-effect adjustment to beginning retained earnings. The Company adopted the provisions of the guidance in the first quarter of 2011. The adoption did not have a material impact on the Company s consolidated financial statements.

In December 2010, the FASB issued an accounting standards update that requires an entity to disclose pro forma revenue and earnings of the combined entity for both the year in which a business combination occurred and the prior year as if the business combination had occurred as of the beginning of prior year only. This guidance became effective prospectively for business combinations occurring in fiscal years beginning after December 15, 2010. The Company adopted the provisions of the guidance in the first quarter of 2011. The adoption did not have a material impact on the Company s consolidated financial statements.

In April 2010, the FASB issued an accounting standards update that provides guidance on the milestone method of revenue recognition for research and development arrangements. This guidance allows an entity to make an accounting policy election to recognize revenue that is contingent upon the achievement of a substantive milestone in its entirety in the period in which the milestone is achieved. This guidance became effective for fiscal years beginning on or after June 15, 2010 and may be applied prospectively to milestones achieved after the adoption date or retrospectively for all periods presented, with earlier application permitted. The Company made an accounting policy election to apply the guidance prospectively beginning in the first quarter of 2011 to recognize revenue in its entirety in the period in which a substantive milestone is achieved. The adoption did not have a material impact on the Company s consolidated financial statements. As of March 31, 2011, the Company has potential future milestone receipts of approximately $473.0 million for the achievement of development, regulatory, and sales milestones in connection with certain collaboration agreements, including $373.0 million related to a development and commercialization agreement that the Company entered into in 2010 with Bristol-Myers Squibb Company that granted Bristol-Myers Squibb Company exclusive worldwide rights to develop, manufacture and commercialize an investigational drug for neuropathic pain. Due to the challenges associated

with developing and obtaining

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

approval for pharmaceutical products, there is substantial uncertainty whether any of the future milestones will be achieved. The Company evaluates whether milestone payments are substantive based on the facts and circumstances associated with each milestone payment in the period it is received.

In October 2009, the FASB issued an accounting standards update that requires an entity to allocate arrangement consideration at the inception of an arrangement to all of its deliverables based on their relative selling prices, eliminates the use of the residual method of allocation, and requires the relative-selling-price method in all circumstances in which an entity recognizes revenue of an arrangement with multiple deliverables. This guidance became effective for revenue arrangements entered into or materially modified in fiscal years beginning on or after June 15, 2010, with earlier application permitted. The Company adopted the provisions of the guidance in the first quarter of 2011. The adoption did not have a material impact on the Company s consolidated financial statements.

Note 2: Acquisitions and Collaborations

Purchase of Distributor s Business in Turkey

On July 1, 2010, the Company terminated its existing distributor agreement in Turkey and completed the purchase from its distributor of all licenses, registrations and other assets related to the selling of the Company s products in Turkey. Additionally, former employees of the distributor who were primarily engaged in the selling and marketing of the Company s products were transferred to the Company on that date. The termination of the existing distributor agreement and purchase of the commercial assets enabled the Company to initiate direct selling operations in Turkey.

In conjunction with the termination of the existing distributor agreement, the Company paid $33.0 million, including a termination fee and related taxes, which is included in selling, general and administrative (SG&A) expenses in the third quarter of 2010. The purchase of the commercial assets was accounted for as a business combination. In connection with the purchase of the assets, the Company paid $6.1 million and is required to pay additional contingent consideration based on specified percentages of revenue in Turkey over the next five years. The estimated fair value of the contingent consideration as of the acquisition date was $36.7 million. The Company recognized goodwill of $31.5 million and intangible assets of $11.3 million based on their estimated fair values at the purchase date. No liabilities were assumed in connection with the purchase. In the first quarter of 2011, the Company made a contingent consideration payment of $2.9 million. As of March 31, 2011, the total estimated fair value of the contingent consideration was $41.6 million, the majority of which was included in Other liabilities.

Serica Acquisition

On January 15, 2010, the Company completed the acquisition of Serica Technologies, Inc. (Serica), a development stage medical device company based in the United States focused on developing biodegradable silk-based scaffolds for use in tissue regeneration for breast reconstruction, for an aggregate purchase price of approximately $63.7 million, net of cash acquired. In connection with the acquisition, the Company acquired assets with a fair value of $96.0 million and assumed liabilities of $32.3 million. The acquisition was funded from the Company s cash and equivalents balances. The Serica acquisition provides the Company with an approved technology that has potential future application in breast augmentation, revision surgeries, as well as potential bariatric applications.

Collaborations

In March 2010, the Company and Serenity Pharmaceuticals, LLC (Serenity) entered into an agreement for the license, development and commercialization of a Phase III investigational drug currently in clinical development for the treatment of nocturia, a common urological disorder in adults characterized by frequent urination at night time. Under the terms of the agreement, the Company receives exclusive worldwide rights to develop, manufacture and commercialize the investigational drug for all potential indications except Primary Nocturnal Enuresis (pediatric bedwetting). In conjunction with the agreement, the Company made an upfront payment to Serenity of $43.0 million in 2010. The terms of the agreement also include potential future development and regulatory milestone payments to Serenity of up to $122.0 million, as well as potential future sales milestone and royalty payments. Because the technology has not yet achieved regulatory approval, the Company recorded the upfront payment of $43.0 million as research and development (R&D) expense in the first quarter of 2010.

In December 2010, the Company and Serenity executed a letter agreement which specified certain terms and conditions governing additional development activities for a new Phase III trial which were not set forth in the original agreement. Under the letter agreement, the Company has agreed to share 50% of the cost of additional development activities. The execution of the letter agreement was a reconsideration event for the Company s variable interest in the collaboration

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

agreement with Serenity, and since the Company is providing a significant amount of the funding for the new Phase III trial, it determined that Serenity had become a variable interest entity (VIE). However, the Company determined that it is not the primary beneficiary of the VIE because it does not possess the power to direct Serenity s research and development activities, which are the activities that most significantly impact Serenity s economic performance. The Company s maximum exposure to loss is the upfront payment of $43.0 million made to Serenity and any shared costs of additional development activities.

On January 28, 2011, the Company entered into a collaboration agreement and a co-promotion agreement with MAP Pharmaceuticals, Inc. (MAP) for the exclusive development and commercialization by the Company and MAP of *Levadex* within the United States to certain headache specialist physicians for the treatment of acute migraine in adults, migraine in adolescents and other indications that may be approved by the parties. *Levadex* is a self-administered, orally inhaled therapy consisting of a proprietary formulation of dihydroergotamine delivered using MAP s proprietary *Temp®* delivery system, which has completed Phase III clinical development for the treatment of acute migraine in adults. MAP currently intends to submit its New Drug Application for *Levadex* to the United States Food and Drug Administration in the first half of 2011. Under the terms of the agreements, the Company made a $60.0 million upfront payment to MAP in February 2011, which was recorded as SG&A expense in the first quarter of 2011. The upfront payment was expensed because *Levadex* has not yet achieved regulatory approval. The terms of the agreements also include up to $97.0 million in additional payments to MAP upon MAP meeting certain development and regulatory milestones.

Note 3: Restructuring Charges and Integration Costs

Discontinued Development of EasyBand

In March 2011, the Company decided to discontinue development of the *EasyBand* Remote Adjustable Gastric Band System (*EasyBand*), a technology that the Company acquired in connection with its 2007 acquisition of EndoArt SA, and close the related research and development facility in Switzerland.

As a result of discontinuing the development of *EasyBand* and the closure of the related research and development facility, in the first quarter of 2011 the Company recorded a pre-tax impairment charge of $16.1 million for the intangible assets associated with the *EasyBand* technology, fixed asset impairment charges of $2.3 million and a gain of $9.4 million from the substantially complete liquidation of the Company s investment in a foreign subsidiary. In addition, the Company recorded $4.6 million of restructuring charges, consisting of $3.0 million of employee severance and other one-time termination benefits for approximately 30 people affected by the facility closure, $1.5 million of contract termination costs and $0.1 million of other related costs.

Other Restructuring Activities and Integration Costs

The Company did not incur any other restructuring charges during the three month period ended March 31, 2011.

Included in the three month period ended March 31, 2010 are $0.1 million of restructuring charges primarily for employee severance and other one-time termination benefits related to the Company s fiscal year 2009 restructuring plan, $0.4 million of restructuring charges primarily for employee severance related to the Serica acquisition and $0.1 million of restructuring charges for an abandoned leased facility related to the Company s fiscal year 2005 restructuring and streamlining of its European operations.

Included in the three month period ended March 31, 2011 are $0.6 million of SG&A expenses related to transaction costs associated with the collaboration and co-promotion agreements with MAP, $0.2 million of SG&A expenses related to integration and transaction costs associated with the purchase of the Company s distributor s business related to the Company s products in Turkey and $0.2 million of SG&A expenses related to transaction costs associated with the announced purchase of the Company s distributor s business related to the Company s products in South Africa.

Included in the three month period ended March 31, 2010 are $0.5 million of SG&A expenses related to integration and transaction costs associated with the Serica acquisition, $0.2 million of SG&A expenses related to transaction costs associated with the purchase of the Company s distributor s business related to the Company s products in Turkey and $0.3 million of SG&A expenses related to transaction costs associated

with the license, development and commercialization agreement with Serenity.

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Note 4: Intangibles and Goodwill

Intangibles

At March 31, 2011 and December 31, 2010, the components of intangibles and certain other related information were as follows:

	March 31, 2011			December 31, 2010		
	Gross Amount	Accumulated Amortization	Weighted Average Amortization Period (in years)	Gross Amount	Accumulated Amortization	Weighted Average Amortization Period (in years)
	(in millions)			(in millions)		
Amortizable Intangible Assets:						
Developed technology	$ 1,117.5	$ (372.5)	13.4	$ 1,129.6	$ (353.2)	13.4
Customer relationships	42.3	(42.3)	3.1	42.3	(42.3)	3.1
Licensing	185.7	(121.8)	9.3	185.6	(116.7)	9.3
Trademarks	27.6	(25.4)	6.3	27.4	(24.2)	6.3
Core technology	184.9	(63.6)	15.2	189.6	(61.5)	15.2
Other	17.2	(2.5)	9.0	17.0	(1.9)	9.1
	1,575.2	(628.1)	12.7	1,591.5	(599.8)	12.7
Unamortizable Intangible Assets:						
In-process research and development	4.3			4.3		
	$ 1,579.5	$ (628.1)		$ 1,595.8	$ (599.8)	

Developed technology consists primarily of current product offerings, primarily breast aesthetics products, obesity intervention products, dermal fillers, skin care products and eye care products acquired in connection with business combinations, asset acquisitions and initial licensing transactions for products previously approved for marketing. Customer relationship assets consist of the estimated value of relationships with customers acquired in connection with the Company s 2006 acquisition of Inamed Corporation (Inamed), primarily in the breast implant market in the United States. Licensing assets consist primarily of capitalized payments to third party licensors related to the achievement of regulatory approvals to commercialize products in specified markets and up-front payments associated with royalty obligations for products that have achieved regulatory approval for marketing. Core technology consists of proprietary technology associated with silicone gel breast implants, gastric bands and intragastric balloon systems acquired in connection with the Inamed acquisition, dermal filler technology acquired in connection with the Company s 2007 acquisition of Groupe Cornéal Laboratoires, and a drug delivery technology acquired in connection with the Company s 2003 acquisition of Oculex Pharmaceuticals, Inc. Other intangible assets consist primarily of acquired product registration rights, distributor relationships, government permits and non-compete agreements. The in-process research and development asset consists of a dermal filler technology that has not yet achieved regulatory approval acquired in connection with the Company s 2010 acquisition of Serica.

In the first quarter of 2011, the Company recorded a pre-tax charge of $16.1 million related to the impairment of the developed technology and core technology associated with *EasyBand* as a result of the discontinued development of the technology.

The following table provides amortization expense by major categories of acquired amortizable intangible assets for the three month periods ended March 31, 2011 and 2010, respectively:

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
Developed technology	$ 22.6	$ 26.6
Customer relationships		0.3
Licensing	5.1	5.8
Trademarks	1.1	1.1
Core technology	3.2	3.1
Other	0.5	0.2
	$ 32.5	$ 37.1

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Amortization expense related to acquired intangible assets generally benefits multiple business functions within the Company, such as the Company s ability to sell, manufacture, research, market and distribute products, compounds and intellectual property. The amount of amortization expense excluded from cost of sales consists primarily of amounts amortized with respect to developed technology and licensing intangible assets.

Estimated amortization expense is $126.1 million for 2011, $120.3 million for 2012, $106.1 million for 2013, $101.1 million for 2014 and $96.0 million for 2015.

Goodwill

Changes in the carrying amount of goodwill by operating segment through March 31, 2011 were as follows:

	Specialty Pharmaceuticals	Medical Devices	Total
		(in millions)	
Balance at December 31, 2010	$ 106.4	$ 1,932.2	$ 2,038.6
Foreign exchange translation effects	0.7	6.0	6.7
Balance at March 31, 2011	$ 107.1	$ 1,938.2	$ 2,045.3

Note 5: Inventories

Components of inventories were:

	March 31, 2011	December 31, 2010
	(in millions)	
Finished products	$ 155.0	$ 148.2
Work in process	26.9	41.1
Raw materials	57.3	40.1
Total	$ 239.2	$ 229.4

At March 31, 2011 and December 31, 2010, approximately $6.9 million and $6.4 million, respectively, of the Company s finished goods inventories, primarily breast implants, were held on consignment at a large number of doctors offices, clinics and hospitals worldwide. The value and quantity at any one location are not significant.

Note 6: Convertible Notes

In 2006, the Company issued its 1.50% Convertible Senior Notes due 2026 (2026 Convertible Notes) for an aggregate principal amount of $750.0 million. The 2026 Convertible Notes are unsecured and pay interest semi-annually on the principal amount of the notes at a rate of 1.50% per annum. The 2026 Convertible Notes mature on April 1, 2026, unless previously redeemed by the Company or earlier converted by the note holders. In the first quarter of 2009, the Company paid $98.3 million to repurchase $100.3 million principal amount of the 2026 Convertible Notes with a carrying value of $92.3 million and a calculated fair value of approximately $97.0 million. The Company separately measures and accounts for the liability and equity components of the 2026 Convertible Notes. As of March 31, 2011, the carrying value of the

liability component is $648.9 million with an effective interest rate of 5.59%. The difference between the carrying value of the liability component and the outstanding principal amount of the 2026 Convertible Notes of $649.7 million is recorded as debt discount and is being amortized to interest expense through the first note holder put date in April 2011.

The 2026 Convertible Notes become convertible into cash and, if applicable, shares of the Company s common stock based on a conversion rate of 15.7904 shares of the Company s common stock per $1,000 principal amount of the 2026 Convertible Notes if the Company s stock price reaches certain specified thresholds or the Company calls the 2026 Convertible Notes for redemption. The Company is permitted to redeem the 2026 Convertible Notes at the principal amount plus accrued interest at any time on or after April 5, 2011.

On March 8, 2011, the Company announced its intention to redeem the 2026 Convertible Notes at the principal amount plus accrued interest on April 5, 2011. Under the terms of the 2026 Convertible Notes, note holders became able to surrender their notes for conversion upon the issuance of the Company s notice of redemption.

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Note 7: Income Taxes

The provision for income taxes is determined using an estimated annual effective tax rate, which is generally less than the U.S. federal statutory rate, primarily because of lower tax rates in certain non-U.S. jurisdictions, R&D tax credits available in the United States, California and other foreign jurisdictions and deductions available in the United States for domestic production activities. The effective tax rate may be subject to fluctuations during the year as new information is obtained, which may affect the assumptions used to estimate the annual effective tax rate, including factors such as the mix of pre-tax earnings in the various tax jurisdictions in which the Company operates, valuation allowances against deferred tax assets, the recognition or derecognition of tax benefits related to uncertain tax positions, expected utilization of R&D tax credits and changes in or the interpretation of tax laws in jurisdictions where the Company conducts business. The Company recognizes deferred tax assets and liabilities for temporary differences between the financial reporting basis and the tax basis of its assets and liabilities along with net operating loss and tax credit carryovers.

The Company records a valuation allowance against its deferred tax assets to reduce the net carrying value to an amount that it believes is more likely than not to be realized. When the Company establishes or reduces the valuation allowance against its deferred tax assets, the provision for income taxes will increase or decrease, respectively, in the period such determination is made. The valuation allowance against deferred tax assets was $23.8 million and $4.3 million as of March 31, 2011 and December 31, 2010, respectively. The increase in the valuation allowance was primarily due to a corresponding increase in a deferred tax asset that the Company determined required a valuation allowance.

The total amount of unrecognized tax benefits was $37.4 million and $32.5 million as of March 31, 2011 and December 31, 2010, respectively. The total amount of unrecognized tax benefits that, if recognized, would affect the effective tax rate was $32.5 million and $27.5 million as of March 31, 2011 and December 31, 2010, respectively. The Company expects that during the next 12 months it is reasonably possible that unrecognized tax benefit liabilities will decrease by approximately $8.0 million to $10.0 million due to the settlement of income tax audits in the United States and certain foreign jurisdictions.

Total interest accrued related to uncertainty in income taxes included in the Company s unaudited condensed consolidated balance sheet was $8.6 million and $8.1 million as of March 31, 2011 and December 31, 2010, respectively.

The Company has not provided for withholding and U.S. taxes for the unremitted earnings of certain non-U.S. subsidiaries because it has currently reinvested these earnings indefinitely in these foreign operations. At December 31, 2010, the Company had approximately $2,109.4 million in unremitted earnings outside the United States for which withholding and U.S. taxes were not provided. Income tax expense would be incurred if these funds were remitted to the United States. It is not practicable to estimate the amount of the deferred tax liability on such unremitted earnings. Upon remittance, certain foreign countries impose withholding taxes that are then available, subject to certain limitations, for use as credits against the Company s U.S. tax liability, if any. The Company annually updates its estimate of unremitted earnings outside the United States after the completion of each fiscal year.

Note 8: Share-Based Compensation

The Company recognizes compensation expense for all share-based awards made to employees and directors. The fair value of share-based awards is estimated at the grant date using the Black-Scholes option-pricing model and the portion that is ultimately expected to vest is recognized as compensation cost over the requisite service period using the straight-line single option method. The fair value of modifications to share-based awards is generally estimated using a lattice model.

The determination of fair value using the Black-Scholes and lattice option-pricing models is affected by the Company s stock price as well as assumptions regarding a number of complex and subjective variables, including expected stock price volatility, risk-free interest rate, expected dividends and projected employee stock option exercise behaviors. The Company currently estimates stock price volatility based upon an equal weighting of the historical average over the expected life of the award and the average implied volatility of at-the-money options traded in the open market. The Company estimates employee stock option exercise behavior based on actual historical exercise activity and assumptions regarding future exercise activity of unexercised, outstanding options.

Share-based compensation expense is recognized only for those awards that are ultimately expected to vest, and the Company has applied an estimated forfeiture rate to unvested awards for the purpose of calculating compensation cost. These estimates will be revised in future periods if actual forfeitures differ from the estimates. Changes in forfeiture estimates impact compensation cost in the period in which the change in estimate occurs.

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

For the three month periods ended March 31, 2011 and 2010, share-based compensation expense was as follows:

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
Cost of sales	$ 1.5	$ 1.1
Selling, general and administrative	13.6	12.9
Research and development	5.7	4.2
Pre-tax share-based compensation expense	20.8	18.2
Income tax benefit	7.4	5.6
Net share-based compensation expense	$ 13.4	$ 12.6

As of March 31, 2011, total compensation cost related to non-vested stock options and restricted stock not yet recognized was approximately $215.9 million, which is expected to be recognized over the next 48 months (38 months on a weighted-average basis). The Company has not capitalized as part of inventory any share-based compensation costs because such costs were negligible as of March 31, 2011.

Note 9: Employee Retirement and Other Benefit Plans

The Company sponsors various qualified defined benefit pension plans covering a substantial portion of its employees. In addition, the Company sponsors two supplemental nonqualified plans covering certain management employees and officers and one retiree health plan covering U.S. retirees and dependents.

Components of net periodic benefit cost for the three month periods ended March 31, 2011 and 2010, respectively, were as follows:

	Three months ended			
	Pension Benefits		Other Postretirement Benefits	
	March 31, 2011	March 31, 2010	March 31, 2011	March 31, 2010
	(in millions)		(in millions)	
Service cost	$ 6.0	$ 5.1	$ 0.6	$ 0.6
Interest cost	10.7	9.8	0.8	0.8
Expected return on plan assets	(11.1)	(11.6)		
Amortization of prior service costs			(0.1)	(0.1)
Recognized net actuarial losses	4.3	2.5	0.2	0.3
Net periodic benefit cost	$ 9.9	$ 5.8	$ 1.5	$ 1.6

In 2011, the Company expects to pay contributions of between $35.0 million and $45.0 million for its U.S. and non-U.S. pension plans and between $1.0 million and $2.0 million for its other postretirement plan.

Note 10: Legal Proceedings

The following supplements and amends the discussion set forth in Note 13 Legal Proceedings in the Company s Annual Report on Form 10-K for the year ended December 31, 2010.

Clayworth v. Allergan, et al.

In August 2004, James Clayworth, R.Ph., doing business as Clayworth Pharmacy, filed a complaint entitled Clayworth v. Allergan, *et al*. in the Superior Court of the State of California for the County of Alameda. The complaint, as amended, named the Company and 12 other defendants and alleged unfair business practices, including a price fixing conspiracy relating to the reimportation of pharmaceuticals from Canada. The complaint sought damages, equitable relief, attorneys fees and costs. In January 2007, the superior court entered a notice of entry of judgment of dismissal against the plaintiffs, dismissing the plaintiffs complaint. On the same date, the plaintiffs filed a notice of appeal with the Court of Appeal of the State of California. In April 2007, the plaintiffs filed an opening brief with the court of appeal. The defendants filed their joint opposition in July 2007, and the plaintiffs filed their reply in August 2007. In May 2008, the court of appeal heard oral arguments and took the matter under submission. In July 2008, the court of appeal affirmed the superior court s ruling, granting the Company s motion for summary judgment. In August 2008, the plaintiffs filed a petition for rehearing with the court of appeal, which the court denied. In September 2008, the plaintiffs filed a petition for review with the

ALLERGAN, INC.

NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Supreme Court of the State of California, which the supreme court granted in November 2008. In February 2009, the plaintiffs filed their opening brief on the merits with the supreme court and defendants filed their answer brief in May 2009. In June 2009, the plaintiffs filed their reply brief on the merits with the supreme court. In May 2010, the supreme court heard oral arguments. In July 2010, the supreme court reversed the court of appeal s judgment and remanded the case to the superior court for further proceedings. In October 2010, plaintiffs filed a challenge to the assignment of this matter to the presiding judge alleging a conflict of interest. In November 2010, plaintiffs challenge was denied. In December 2010, plaintiffs filed a petition for writ of mandate in the Court of Appeal of the State of California seeking to overturn the order denying their challenge. In December 2010, the court of appeal denied the petition. In December 2010, plaintiffs filed a petition for review with the Supreme Court of the State of California. In January 2011, the court set trial for August 1, 2011. In February 2011, the supreme court denied plaintiffs petition for review. In March 2011, the court entered judgment in favor of defendants pursuant to orders granting motions for summary judgment. In April 2011, plaintiffs filed a notice of appeal to the Court of Appeal of the State of California.

Kramer et al. v. Allergan, Inc.

In July 2008, a complaint entitled Kramer, Bryant, Spears, Doolittle, Clark, Whidden, Powell, Moore, Hennessey, Sody, Breeding, Downey, Underwood-Boswell, Reed-Momot, Purdon & Hahn v. Allergan, Inc. was filed in the Superior Court for the State of California for the County of Orange. The complaint makes allegations against the Company relating to *Botox*® and *Botox*® Cosmetic including failure to warn, manufacturing defects, negligence, breach of implied and express warranties, deceit by concealment and negligent misrepresentation and seeks damages, attorneys fees and costs. In 2009, the plaintiffs Hennessey, Hahn, Underwood-Boswell, Purdon, Moore, Clark, Reed-Momot and Whidden were dismissed without prejudice. In October 2009, the Company filed a motion for summary judgment against plaintiff Spears, which the court denied in December 2009. The trial related to plaintiff Spears began in January 2010. In March 2010, the jury returned a verdict in the Company s favor and the court entered a judgment on the special verdict. In April 2010, plaintiff Spears filed a motion for a new trial which the court denied in May 2010. In June 2010, the Company and plaintiff Spears entered into a settlement agreement under which the Company agreed to waive costs in exchange for plaintiff Spears agreeing not to appeal the judgment. In September 2010, the trial related to plaintiff Bryant began and the Company subsequently entered into a settlement agreement with plaintiff Bryant. In April 2011, the court set the trial related to plaintiff Doolittle for November 7, 2011 or, in the alternative, for January 17, 2012.

Government Investigations

In June 2010, the Company received service of process of a Subpoena from the U.S. Securities and Exchange Commission (SEC). The subpoena requests the production of documents relating to the Company s affiliation with Acadia Pharmaceuticals, Inc., or Acadia, and the Company s sale of Acadia securities. In September 2010, the Company produced documents responsive to the Subpoena. In January and March 2011, the SEC issued additional Subpoenas seeking further information, which has been provided.

Stockholder Derivative Litigation

In November 2010, the Company received a demand for inspection of books and records from U.F.C.W. Local 1776 & Participating Employers Pension Fund (U.F.C.W.). In November 2010, U.F.C.W. filed a motion to intervene in the Louisiana Municipal Police Employees Retirement System action, which was denied by the court in January 2011. In February 2011, U.F.C.W. filed a complaint to compel inspection of books and records in the Court of Chancery of the State of Delaware. In March 2011, the Company filed an answer to the complaint and the court scheduled the final hearing for April 27, 2011. In April 2011, the court ordered that the Company produce a limited number of documents to the court for in camera inspection.

In September 2010, Pompano Beach Police & Firefighters Retirement System and Western Washington Laborers-Employers Pension Trust filed a stockholder derivative complaint against the Company s then-current Board of Directors and Allergan, Inc. in the U.S. District Court for the Central District of California. The complaint alleges violations of federal securities laws, breaches of fiduciary duties, abuse of control, gross mismanagement, and corporate waste and seeks, among other things, damages, corporate governance reforms, attorneys fees, and costs. In September 2010, plaintiffs filed a motion for consolidation with the Himmel and Rosenbloom actions, which the court granted in October 2010. In November 2010, the plaintiffs filed their consolidated complaint. In December 2010, the Company filed a motion to stay the consolidated action in favor of the Louisiana Municipal Police Employees Retirement System action. In December 2010, the Company and the individual defendants filed motions to dismiss the consolidated complaint. In March 2011, the court denied the motion to stay the consolidated action and

the Company filed a motion for partial stay of the consolidated action in favor of the Louisiana Municipal Police Employees Retirement System action. In April 2011, the court granted the motions to dismiss the consolidated complaint with leave to amend.

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The Company is involved in various other lawsuits and claims arising in the ordinary course of business. These other matters are, in the opinion of management, immaterial both individually and in the aggregate with respect to the Company s consolidated financial position, liquidity or results of operations. Because of the uncertainties related to the incurrence, amount and range of loss on any pending litigation, investigation, inquiry or claim, management is currently unable to predict the ultimate outcome of any litigation, investigation, inquiry or claim, determine whether a liability has been incurred or make an estimate of the reasonably possible liability that could result from an unfavorable outcome. The Company believes however, that the liability, if any, resulting from the aggregate amount of uninsured damages for any outstanding litigation, investigation or claim will not have a material adverse effect on the Company s consolidated financial position, liquidity or results of operations. However, an adverse ruling in a patent infringement lawsuit involving the Company could materially affect the Company s ability to sell one or more of its products or could result in additional competition. In view of the unpredictable nature of such matters, the Company cannot provide any assurances regarding the outcome of any litigation, investigation, inquiry or claim to which the Company is a party or the impact on the Company of an adverse ruling in such matters.

Note 11: Contingencies

In 2009, the Company established a reserve for a contingent liability associated with regulation changes resulting from a final rule issued by the U.S. Department of Defense (DoD) that placed retroactive and prospective pricing limits on certain branded pharmaceuticals under the TRICARE Retail Pharmacy Program, even though such branded pharmaceuticals have not historically been subject to a contract with the Company. The Company is currently in negotiations with the DoD to seek a waiver of retroactive rebates. As of March 31, 2011, the reserve for the contingent liability is $12.4 million and is included in Other accrued expenses.

In the third quarter of 2009, the Company entered into a co-promotion agreement with Quintiles Transnational Corp. (Quintiles), under which Quintiles co-promoted *Sanctura XR*®, *Latisse*® and *Aczone*®, generally targeting primary care physicians. Due to significantly lower than anticipated performance under the agreement, the Company terminated this co-promotion agreement in the third quarter of 2010. The Company estimates it will be required to pay between approximately $15.0 million and $25.0 million in costs in connection with the termination of the co-promotion agreement. As of March 31, 2011, the Company is carrying a recorded reserve for this contingent liability within the range specified above, which is included in Other accrued expenses.

Note 12: Guarantees

The Company s Amended and Restated Certificate of Incorporation provides that the Company will indemnify, to the fullest extent permitted by the Delaware General Corporation Law, each person that is involved in or is, or is threatened to be, made a party to any action, suit or proceeding by reason of the fact that he or she, or a person of whom he or she is the legal representative, is or was a director or officer of the Company or was serving at the request of the Company as a director, officer, employee or agent of another corporation or of a partnership, joint venture, trust or other enterprise. The Company has also entered into contractual indemnity agreements with each of its directors and executive officers pursuant to which, among other things, the Company has agreed to indemnify such directors and executive officers against any payments they are required to make as a result of a claim brought against such executive officer or director in such capacity, excluding claims (i) relating to the action or inaction of a director or executive officer that resulted in such director or executive officer gaining illegal personal profit or advantage, (ii) for an accounting of profits made from the purchase or sale of securities of the Company within the meaning of Section 16(b) of the Securities Exchange Act of 1934, as amended, or similar provisions of any state law or (iii) that are based upon or arise out of such director s or executive officer s knowingly fraudulent, deliberately dishonest or willful misconduct. The maximum potential amount of future payments that the Company could be required to make under these indemnification provisions is unlimited. However, the Company has purchased directors and officers liability insurance policies intended to reduce the Company s monetary exposure and to enable the Company to recover a portion of any future amounts paid. The Company has not previously paid any material amounts to defend lawsuits or settle claims as a result of these indemnification provisions, but makes no assurance that such amounts will not be paid in the future. The Company currently believes the estimated fair value of these indemnification arrangements is minimal.

The Company customarily agrees in the ordinary course of its business to indemnification provisions in agreements with clinical trials investigators in its drug, biologics and medical device development programs, in sponsored research agreements with academic and not-for-profit institutions, in various comparable agreements involving parties performing services for the Company in the ordinary course of business, and in its real estate leases. The Company also customarily agrees to certain indemnification provisions in its discovery and development collaboration agreements. With respect to the Company s clinical trials and sponsored research agreements, these indemnification

provisions typically apply to any claim asserted against the investigator or the investigator s institution relating to personal injury or property damage, violations of

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law or certain breaches of the Company s contractual obligations arising out of the research or clinical testing of the Company s products, compounds or drug candidates. With respect to real estate lease agreements, the indemnification provisions typically apply to claims asserted against the landlord relating to personal injury or property damage caused by the Company, to violations of law by the Company or to certain breaches of the Company s contractual obligations. The indemnification provisions appearing in the Company s collaboration agreements are similar, but in addition provide some limited indemnification for the collaborator in the event of third party claims alleging infringement of intellectual property rights. In each of the above cases, the terms of these indemnification provisions generally survive the termination of the agreement. The maximum potential amount of future payments that the Company could be required to make under these provisions is generally unlimited. The Company has purchased insurance policies covering personal injury, property damage and general liability intended to reduce the Company s exposure for indemnification and to enable the Company to recover a portion of any future amounts paid. The Company has not previously paid any material amounts to defend lawsuits or settle claims as a result of these indemnification provisions. As a result, the Company believes the estimated fair value of these indemnification arrangements is minimal.

Note 13: Product Warranties

The Company provides warranty programs for breast implant sales primarily in the United States, Europe and certain other countries. Management estimates the amount of potential future claims from these warranty programs based on actuarial analyses. Expected future obligations are determined based on the history of product shipments and claims and are discounted to a current value. The liability is included in both current and long-term liabilities in the Company s consolidated balance sheets. The U.S. programs include the *ConfidencePlus*® and *ConfidencePlus*® Premier warranty programs. The *ConfidencePlus*® program currently provides lifetime product replacement, $1,200 of financial assistance for surgical procedures within ten years of implantation and contralateral implant replacement. The *ConfidencePlus*® Premier program, which normally requires a low additional enrollment fee, generally provides lifetime product replacement, $2,400 of financial assistance for saline breast implants and $3,500 of financial assistance for silicone gel breast implants for surgical procedures within ten years of implantation and contralateral implant replacement. The enrollment fee is deferred and recognized as income over the ten year warranty period for financial assistance. The warranty programs in non-U.S. markets have similar terms and conditions to the U.S. programs. The Company does not warrant any level of aesthetic result and, as required by government regulation, makes extensive disclosures concerning the risks of the use of its products and breast implant surgery. Changes to actual warranty claims incurred and interest rates could have a material impact on the actuarial analysis and the Company s estimated liabilities. A large majority of the product warranty liability arises from the U.S. warranty programs. The Company does not currently offer any similar warranty program on any other product.

The following table provides a reconciliation of the change in estimated product warranty liabilities through March 31, 2011:

		(in millions)
Balance at December 31, 2010	$	30.1
Provision for warranties issued during the period		0.9
Settlements made during the period		(0.6)
Decreases in warranty estimates		(0.1)
Balance at March 31, 2011	$	30.3
Current portion	$	6.6
Non-current portion		23.7
Total	$	30.3

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Note 14: Earnings Per Share

The table below presents the computation of basic and diluted earnings per share:

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions, except per share amounts)	
Net earnings attributable to Allergan, Inc.	$ 158.3	$ 167.9
Weighted average number of shares outstanding	304.5	303.5
Net shares assumed issued using the treasury stock method for options and non-vested equity shares and share units outstanding during each period based on average market price	5.1	3.6
Dilutive effect of assumed conversion of convertible notes outstanding	1.2	
Diluted shares	310.8	307.1
Earnings per share attributable to Allergan, Inc. stockholders:		
Basic	$ 0.52	$ 0.55
Diluted	$ 0.51	$ 0.55

For the three month periods ended March 31, 2011 and 2010, options to purchase 4.8 million and 11.2 million shares of common stock at exercise prices ranging from $62.71 to $75.58 and $47.10 to $65.63 per share, respectively, were outstanding but were not included in the computation of diluted earnings per share because the effect from the assumed exercise of these options calculated under the treasury stock method would be anti-dilutive. There were no potentially diluted common shares related to the Company s 2026 Convertible Notes for the three month period ended March 31, 2010, as the Company s average stock price for the period was less than the conversion price of the notes.

Note 15: Comprehensive Income (Loss)

The following table summarizes the components of comprehensive income (loss) for the three month periods ended March 31, 2011 and 2010:

	Three months ended					
	March 31, 2011			March 31, 2010		
	Before Tax Amount	Tax (Expense) or Benefit	Net-of-Tax Amount	Before Tax Amount	Tax (Expense) or Benefit	Net-of-Tax Amount
	(in millions)					
Foreign currency translation adjustments	$ 22.0	$	$ 22.0	$ (19.2)	$	$ (19.2)
Amortization of deferred holding gains on derivatives designated as cash flow hedges	(0.3)	0.1	(0.2)	(0.3)	0.1	(0.2)

Other comprehensive income (loss)	$ 21.7	$ 0.1	21.8	$ (19.5)	$ 0.1	(19.4)
Net earnings			158.8			169.0
Total comprehensive income			180.6			149.6
Comprehensive income attributable to noncontrolling interest			1.1			1.8
Comprehensive income attributable to Allergan, Inc.			$ 179.5			$ 147.8

Note 16: Financial Instruments

In the normal course of business, operations of the Company are exposed to risks associated with fluctuations in interest rates and foreign currency exchange rates. The Company addresses these risks through controlled risk management that includes the use of derivative financial instruments to economically hedge or reduce these exposures. The Company does not enter into derivative financial instruments for trading or speculative purposes.

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The Company has not experienced any losses to date on its derivative financial instruments due to counterparty credit risk.

To ensure the adequacy and effectiveness of its interest rate and foreign exchange hedge positions, the Company continually monitors its interest rate swap positions and foreign exchange forward and option positions both on a stand-alone basis and in conjunction with its underlying interest rate and foreign currency exposures, from an accounting and economic perspective.

However, given the inherent limitations of forecasting and the anticipatory nature of the exposures intended to be hedged, the Company cannot assure that such programs will offset more than a portion of the adverse financial impact resulting from unfavorable movements in either interest or foreign exchange rates. In addition, the timing of the accounting for recognition of gains and losses related to mark-to-market instruments for any given period may not coincide with the timing of gains and losses related to the underlying economic exposures and, therefore, may adversely affect the Company s consolidated operating results and financial position.

Interest Rate Risk Management

The Company s interest income and expense is more sensitive to fluctuations in the general level of U.S. interest rates than to changes in rates in other markets. Changes in U.S. interest rates affect the interest earned on cash and equivalents and short-term investments and interest expense on debt, as well as costs associated with foreign currency contracts.

On January 31, 2007, the Company entered into a nine-year, two-month interest rate swap with a $300.0 million notional amount with semi-annual settlements and quarterly interest rate reset dates. The swap receives interest at a fixed rate of 5.75% and pays interest at a variable interest rate equal to 3-month LIBOR plus 0.368%, and effectively converts $300.0 million of the Company s $800.0 million in aggregate principal amount of 5.75% Senior Notes due 2016 (2016 Notes) to a variable interest rate. Based on the structure of the hedging relationship, the hedge meets the criteria for using the short-cut method for a fair value hedge. The investment in the derivative and the related long-term debt are recorded at fair value. At March 31, 2011 and December 31, 2010, the Company recognized in its consolidated balance sheets an asset reported in Investments and other assets and a corresponding increase in Long-term debt associated with the fair value of the derivative of $37.5 million and $42.3 million, respectively. The differential to be paid or received as interest rates change is accrued and recognized as an adjustment of interest expense related to the 2016 Notes. During the three month periods ended March 31, 2011 and 2010, the Company recognized $3.8 million, respectively, as a reduction of interest expense due to the differential to be received.

In February 2006, the Company entered into interest rate swap contracts based on 3-month LIBOR with an aggregate notional amount of $800.0 million, a swap period of 10 years and a starting swap rate of 5.198%. The Company entered into these swap contracts as a cash flow hedge to effectively fix the future interest rate for the 2016 Notes. In April 2006, the Company terminated the interest rate swap contracts and received approximately $13.0 million. The total gain was recorded to accumulated other comprehensive loss and is being amortized as a reduction to interest expense over a 10 year period to match the term of the 2016 Notes. During the three month periods ended March 31, 2011 and 2010, the Company recognized $0.3 million, respectively, as a reduction of interest expense due to the amortization of deferred holding gains on derivatives designated as cash flow hedges. These amounts were reclassified from accumulated other comprehensive loss. As of March 31, 2011, the remaining unrecognized gain of $6.6 million ($3.9 million, net of tax) is recorded as a component of accumulated other comprehensive loss. The Company expects to reclassify an estimated pre-tax amount of $1.3 million from accumulated other comprehensive loss as a reduction in interest expense during fiscal year 2011 due to the amortization of deferred holding gains on derivatives designated as cash flow hedges.

No portion of amounts recognized from contracts designated as cash flow hedges was considered to be ineffective during the three month periods ended March 31, 2011 and 2010, respectively.

Foreign Exchange Risk Management

Overall, the Company is a net recipient of currencies other than the U.S. dollar and, as such, benefits from a weaker dollar and is adversely affected by a stronger dollar relative to major currencies worldwide. Accordingly, changes in exchange rates, and in particular a strengthening of the U.S. dollar, may negatively affect the Company s consolidated revenues or operating costs and expenses as expressed in U.S. dollars.

From time to time, the Company enters into foreign currency option and forward contracts to reduce earnings and cash flow volatility associated with foreign exchange rate changes to allow management to focus its attention on its core business issues. Accordingly, the Company enters into various contracts which change in value as foreign exchange rates change to

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economically offset the effect of changes in the value of foreign currency assets and liabilities, commitments and anticipated foreign currency denominated sales and operating expenses. The Company enters into foreign currency option and forward contracts in amounts between minimum and maximum anticipated foreign exchange exposures, generally for periods not to exceed 18 months. The Company does not designate these derivative instruments as accounting hedges.

The Company uses foreign currency option contracts, which provide for the sale or purchase of foreign currencies to offset foreign currency exposures expected to arise in the normal course of the Company s business. While these instruments are subject to fluctuations in value, such fluctuations are anticipated to offset changes in the value of the underlying exposures.

Probable but not firmly committed transactions are comprised of sales of products and purchases of raw material in currencies other than the U.S. dollar. A majority of these sales are made through the Company s subsidiaries in Europe, Asia Pacific, Canada and Brazil. The Company purchases foreign exchange option contracts to economically hedge the currency exchange risks associated with these probable but not firmly committed transactions. The duration of foreign exchange hedging instruments, whether for firmly committed transactions or for probable but not firmly committed transactions, generally does not exceed 18 months.

All of the Company s outstanding foreign currency option contracts are entered into to reduce the volatility of earnings generated in currencies other than the U.S. dollar, primarily earnings denominated in the Canadian dollar, Mexican peso, Australian dollar, Brazilian real, euro, Korean won and Turkish lira. Current changes in the fair value of open foreign currency option contracts and any realized gains (losses) on settled contracts are recorded through earnings as Other, net in the accompanying unaudited condensed consolidated statements of earnings. During the three month periods ended March 31, 2011 and 2010, the Company recognized realized gains on settled foreign currency option contracts of $0.5 million and $2.0 million, respectively, and net unrealized losses on open foreign currency option contracts of $6.9 million and $0.7 million, respectively. The premium costs of purchased foreign exchange option contracts are recorded in Other current assets and amortized to Other, net over the life of the options.

All of the Company s outstanding foreign exchange forward contracts are entered into to offset the change in value of certain intercompany receivables or payables that are subject to fluctuations in foreign currency exchange rates. The realized and unrealized gains and losses from foreign currency forward contracts and the revaluation of the foreign denominated intercompany receivables or payables are recorded through Other, net in the accompanying unaudited condensed consolidated statements of earnings. During the three month periods ended March 31, 2011 and 2010, the Company recognized total realized and unrealized gains from foreign exchange forward contracts of $1.7 million and $0.7 million, respectively.

The fair value of outstanding foreign exchange option and forward contracts, collectively referred to as foreign currency derivative financial instruments, are recorded in Other current assets and Accounts payable. At March 31, 2011 and December 31, 2010, foreign currency derivative assets associated with the foreign exchange option contracts of $4.6 million and $10.4 million, respectively, were included in Other current assets. At March 31, 2011 and December 31, 2010, net foreign currency derivative liabilities associated with the foreign exchange forward contracts of $0.2 million and $0.7 million, respectively, were included in Accounts payable.

At March 31, 2011 and December 31, 2010, the notional principal and fair value of the Company s outstanding foreign currency derivative financial instruments were as follows:

	March 31, 2011		December 31, 2010	
	Notional Principal	Fair Value	Notional Principal	Fair Value
	(in millions)			
Foreign currency forward exchange contracts (Receive U.S. dollar/pay foreign currency)	$ 36.3	$ (0.9)	$ 25.6	$ (0.9)

Foreign currency forward exchange contracts				
(Pay U.S. dollar/receive foreign currency)	41.7	0.7	39.9	0.2
Foreign currency sold put options	344.4	4.6	346.4	10.4

The notional principal amounts provide one measure of the transaction volume outstanding as of March 31, 2011 and December 31, 2010, and do not represent the amount of the Company s exposure to market loss. The estimates of fair value are based on applicable and commonly used pricing models using prevailing financial market information as of March 31, 2011 and December 31, 2010. The amounts ultimately realized upon settlement of these financial instruments, together with the gains and losses on the underlying exposures, will depend on actual market conditions during the remaining life of the instruments.

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NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Other Financial Instruments

At March 31, 2011 and December 31, 2010, the Company s other financial instruments included cash and equivalents, short-term investments, trade receivables, equity investments, accounts payable and borrowings. The carrying amount of cash and equivalents, short-term investments, trade receivables and accounts payable approximates fair value due to the short-term maturities of these instruments. The fair value of non-marketable equity investments which represent investments in start-up technology companies or partnerships that invest in start-up technology companies, are estimated based on the fair value and other information provided by these ventures. The fair value of notes payable, convertible notes and long-term debt are estimated based on quoted market prices and interest rates.

The carrying amount and estimated fair value of the Company s other financial instruments at March 31, 2011 and December 31, 2010 were as follows:

	March 31, 2011		December 31, 2010	
	Carrying Amount	Fair Value	Carrying Amount	Fair Value
	(in millions)			
Cash and equivalents	$ 2,530.8	$ 2,530.8	$ 1,991.2	$ 1,991.2
Short-term investments	249.7	249.7	749.1	749.1
Non-current non-marketable equity investments	7.7	7.7	7.7	7.7
Notes payable	36.3	36.3	28.1	28.1
Convertible notes	648.9	649.9	642.5	651.1
Long-term debt	1,529.5	1,589.3	1,534.2	1,612.3

Concentration of Credit Risk

Financial instruments that potentially subject the Company to credit risk principally consist of trade receivables. Wholesale distributors, major retail chains and managed care organizations account for a substantial portion of trade receivables. This risk is limited due to the number of customers comprising the Company s customer base, and their geographic dispersion. At March 31, 2011, no single customer represented more than 10% of trade receivables, net. Ongoing credit evaluations of customers financial condition are performed and, generally, no collateral is required. The Company has purchased an insurance policy intended to reduce the Company s exposure to potential credit risks associated with certain U.S. customers. To date, no claims have been made against the insurance policy. The Company maintains reserves for potential credit losses and such losses, in the aggregate, have not exceeded management s estimates.

Note 17: Fair Value Measurements

The Company measures fair value based on the prices 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. Fair value measurements are based on a three-tier hierarchy that prioritizes the inputs used to measure fair value. These tiers include: Level 1, defined as observable inputs such as quoted prices in active markets; Level 2, defined as inputs other than quoted prices in active markets that are either directly or indirectly observable; and Level 3, defined as unobservable inputs for which little or no market data exists, therefore requiring an entity to develop its own assumptions.

Assets and Liabilities Measured at Fair Value on a Recurring Basis

As of March 31, 2011, the Company has certain assets and liabilities that are required to be measured at fair value on a recurring basis. These include cash equivalents, short-term investments, foreign exchange derivatives, the $300.0 million notional amount interest rate swap and contingent consideration. These assets and liabilities are classified in the table below in one of the three categories of the fair value hierarchy described above.

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ALLERGAN, INC.

NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

	Total	Level 1	Level 2	Level 3
		(in millions)		
Assets				
Commercial paper	$ 1,159.4	$	$ 1,159.4	$
Foreign time deposits	227.5		227.5	
Other cash equivalents	1,272.1		1,272.1	
Foreign exchange derivative assets	4.6		4.6	
Interest rate swap derivative asset	37.5		37.5	
	$ 2,701.1	$	$ 2,701.1	$
Liabilities				
Foreign exchange derivative liabilities	$ 0.2	$	$ 0.2	$
Interest rate swap derivative liability	37.5		37.5	
Contingent consideration liability	41.6			41.6
	$ 79.3	$	$ 37.7	$ 41.6

Cash equivalents consist of commercial paper, foreign time deposits and other cash equivalents. Short-term investments consist of commercial paper. Cash equivalents and short-term investments are valued at cost, which approximates fair value due to the short-term maturities of these instruments. Foreign currency derivative assets and liabilities are valued using quoted forward foreign exchange prices and option volatility at the reporting date. The interest rate swap derivative asset and liability are valued using LIBOR yield curves at the reporting date. The Company believes the fair values assigned to its derivative instruments as of March 31, 2011 are based upon reasonable estimates and assumptions.

The contingent consideration liability represents future amounts the Company will be required to pay in conjunction with the 2010 purchase of commercial assets from a distributor in Turkey that was accounted for as a business combination. The ultimate amount of future payments is based on specified percentages of the Company s revenues in Turkey over the next five years. The Company estimates the fair value of the contingent liability using the income approach, which involves forecasting estimated future net cash flows and discounting the net cash flows to their present value using a risk-adjusted rate of return. As of the acquisition date, the fair value of the liability was estimated to be $36.7 million. As of March 31, 2011 and December 31, 2010, the total estimated fair value of the contingent consideration was $41.6 million and $44.5 million, respectively. In the first quarter of 2011, the Company made a contingent consideration payment of $2.9 million.

Note 18: Business Segment Information

The Company operates its business on the basis of two reportable segments specialty pharmaceuticals and medical devices. The specialty pharmaceuticals segment produces a broad range of pharmaceutical products, including: ophthalmic products for dry eye, glaucoma, retinal diseases and ocular surface disease; *Botox*® for certain therapeutic and aesthetic indications; skin care products for acne, psoriasis, eyelash growth and other prescription and over-the-counter skin care products; and urologics products. The medical devices segment produces a broad range of medical devices, including: breast implants for augmentation, revision and reconstructive surgery; obesity intervention products, including the *Lap-Band*® System and the *Orbera* Intragastric Balloon System; and facial aesthetics products. The Company provides global marketing strategy teams to ensure development and execution of a consistent marketing strategy for its products in all geographic regions that share similar distribution channels and customers.

The Company evaluates segment performance on a revenue and operating income basis exclusive of general and administrative expenses and other indirect costs, legal settlement expenses, intangible asset impairment and related costs, restructuring charges, in-process research and development expenses, amortization of certain identifiable intangible assets related to business combinations and asset acquisitions and related capitalized licensing costs and certain other adjustments, which are not allocated to the Company s segments for performance assessment by the Company s chief operating decision maker. Other adjustments excluded from the Company s segments for performance assessment represent income or expenses that do not reflect, according to established Company-defined criteria, operating income or expenses associated with the Company s core business activities. Because operating segments are generally defined by the products they design and sell, they do not make

sales to each other. The Company does not discretely allocate assets to its operating segments, nor does the Company s chief operating decision maker evaluate operating segments using discrete asset information.

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ALLERGAN, INC.

NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Operating Segments

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
Product net sales:		
Specialty pharmaceuticals	$ 1,028.4	$ 907.3
Medical devices	224.4	198.5
Total product net sales	1,252.8	1,105.8
Other corporate and indirect revenues	18.4	48.9
Total revenues	$ 1,271.2	$ 1,154.7
Operating income:		
Specialty pharmaceuticals	$ 384.2	$ 311.9
Medical devices	67.5	67.1
Total segments	451.7	379.0
General and administrative expenses, other indirect costs and other adjustments	156.9	96.7
Amortization of acquired intangible assets (a)	26.6	31.4
Intangible asset impairment	16.1	
Restructuring charges	4.6	0.6
Total operating income	$ 247.5	$ 250.3

(a) Represents amortization of certain identifiable intangible assets related to business combinations and asset acquisitions and related capitalized licensing costs, as applicable.

Product net sales for the Company s various global product portfolios are presented below. The Company s principal markets are the United States, Europe, Latin America and Asia Pacific. The U.S. information is presented separately as it is the Company s headquarters country. U.S. sales represented 60.9% and 62.5% of the Company s total consolidated product net sales for the three month periods ended March 31, 2011 and 2010, respectively.

Sales to two customers in the Company s specialty pharmaceuticals segment each generated over 10% of the Company s total consolidated product net sales. Sales to Cardinal Health, Inc. for the three month periods ended March 31, 2011 and 2010 were 14.7% and 12.1%, respectively, of the Company s total consolidated product net sales. Sales to McKesson Drug Company for the three month periods ended March 31, 2011 and 2010 were 14.4% and 14.0%, respectively, of the Company s total consolidated product net sales. No other country or single customer generates over 10% of the Company s total consolidated product net sales. Net sales for the Europe region also include sales to customers in Africa and the Middle East, and net sales in the Asia Pacific region include sales to customers in Australia and New Zealand.

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ALLERGAN, INC.

NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

Product Net Sales by Product Line

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
Specialty Pharmaceuticals:		
Eye Care Pharmaceuticals	$ 591.9	$ 512.0
Botox®/Neuromodulators	364.5	331.0
Skin Care	58.7	50.6
Urologics	13.3	13.7
Total Specialty Pharmaceuticals	1,028.4	907.3
Medical Devices:		
Breast Aesthetics	84.1	77.9
Obesity Intervention	52.1	61.2
Facial Aesthetics	88.2	59.4
Total Medical Devices	224.4	198.5
Total product net sales	$ 1,252.8	$ 1,105.8

Geographic Information

Product Net Sales

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
United States	$ 762.7	$ 690.8
Europe	260.0	224.9
Latin America	84.4	63.9
Asia Pacific	92.2	78.6
Other	53.5	47.6
Total product net sales	$ 1,252.8	$ 1,105.8

Long-Lived Assets

	March 31, 2011	December 31, 2010
	(in millions)	
United States	$ 3,194.4	$ 3,222.4
Europe	539.2	563.1
Latin America	64.0	65.0

Asia Pacific	56.2	56.3
Other	3.5	3.7
Total	$ 3,857.3	$ 3,910.5

Note 19: Subsequent Events

On March 8, 2011, the Company announced its intention to redeem the 2026 Convertible Notes at the principal amount plus accrued interest on April 5, 2011. Most note holders have elected to exercise the conversion feature of the 2026 Convertible Notes prior to redemption. Pursuant to the terms of the 2026 Convertible Notes, the Company has elected to pay the full conversion value in cash. The conversion value of a note is based on an average of the daily closing price of the Company s stock over an averaging period that commences after the Company receives a conversion notice from a note holder. The conversion value of the 2026 Convertible Notes will be paid to note holders at the end of the applicable averaging periods in April and May 2011 and the aggregate amount of such conversion value is expected to be approximately $800.0 million. The Company redeemed the notes that were not converted for $8.6 million, representing the aggregate principal amount plus accrued interest, on April 5, 2011.

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ALLERGAN, INC.

NOTES TO UNAUDITED CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (Continued)

In May 2011, the Company announced that effective July 1, 2011 the Company will establish direct operations in South Africa by acquiring the Allergan-related parts of Genop Healthcare s business and assume responsibility for promotion, marketing and distribution of all Allergan products in South Africa. The acquisition will be accounted for as a business combination, the terms of which are not material.

On May 4, 2011, the Company announced a license agreement with Molecular Partners AG pursuant to which the Company obtains exclusive global rights in the field of ophthalmology for MP0112, a Phase II proprietary therapeutic *DARPin*® protein targeting vascular endothelial growth factor receptors under investigation for the treatment of retinal diseases. Under the terms of the agreement, the Company will make a $45.0 million upfront payment to Molecular Partners AG and potential future milestone and royalty payments.

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ALLERGAN, INC.

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

This financial review presents our operating results for the three month periods ended March 31, 2011 and 2010, and our financial condition at March 31, 2011. The following discussion contains forward-looking statements which are subject to known and unknown risks, uncertainties and other factors that may cause our actual results to differ materially from those expressed or implied by such forward-looking statements. We discuss such risks, uncertainties and other factors throughout this report and specifically under the caption Risk Factors in Part II, Item 1A below. The following review should be read in connection with the information presented in our unaudited condensed consolidated financial statements and related notes for the three month period ended March 31, 2011 included in this report and our audited consolidated financial statements and related notes for the year ended December 31, 2010 included in our 2010 Annual Report on Form 10-K filed with the Securities and Exchange Commission.

Critical Accounting Policies, Estimates and Assumptions

The preparation and presentation of financial statements in conformity with accounting principles generally accepted in the United States, or GAAP, requires us to establish policies and to make estimates and assumptions that affect the amounts reported in our consolidated financial statements. In our judgment, the accounting policies, estimates and assumptions described below have the greatest potential impact on our consolidated financial statements. Accounting assumptions and estimates are inherently uncertain and actual results may differ materially from our estimates.

Revenue Recognition

We recognize revenue from product sales when goods are shipped and title and risk of loss transfer to our customers. A substantial portion of our revenue is generated by the sale of specialty pharmaceutical products (primarily eye care pharmaceuticals, skin care and urologics products) to wholesalers within the United States, and we have a policy to attempt to maintain average U.S. wholesaler inventory levels at an amount less than eight weeks of our net sales. A portion of our revenue is generated from consigned inventory of breast implants maintained at physician, hospital and clinic locations. These customers are contractually obligated to maintain a specific level of inventory and to notify us upon the use of consigned inventory. Revenue for consigned inventory is recognized at the time we are notified by the customer that the product has been used. Notification is usually through the replenishing of the inventory, and we periodically review consignment inventories to confirm the accuracy of customer reporting.

We generally offer cash discounts to customers for the early payment of receivables. Those discounts are recorded as a reduction of revenue and accounts receivable in the same period that the related sale is recorded. The amounts reserved for cash discounts were $3.9 million and $4.4 million at March 31, 2011 and December 31, 2010, respectively. Provisions for cash discounts deducted from consolidated sales in the first quarter of 2011 and 2010 were $14.3 million and $12.4 million, respectively.

We permit returns of product from most product lines by any class of customer if such product is returned in a timely manner, in good condition and from normal distribution channels. Return policies in certain international markets and for certain medical device products, primarily breast implants, provide for more stringent guidelines in accordance with the terms of contractual agreements with customers. Our estimates for sales returns are based upon the historical patterns of product returns matched against sales, and management s evaluation of specific factors that may increase the risk of product returns. The amount of allowances for sales returns recognized in our consolidated balance sheets at March 31, 2011 and December 31, 2010 were $58.7 million and $52.3 million, respectively, and are recorded in Other accrued expenses and Trade receivables, net in our consolidated balance sheets. Provisions for sales returns deducted from consolidated sales were $103.9 million and $87.1 million in the first quarter of 2011 and 2010, respectively. The increases in the amount of allowances for sales returns at March 31, 2011 compared to December 31, 2010 and the provisions for sales returns in the first quarter of 2011 compared to the first quarter of 2010 are primarily due to increased sales returns related to breast implant products, principally due to increased product sales volume, and an increase in estimated product return rates for our skin care products. Historical allowances for cash discounts and product returns have been consistent with the amounts reserved or accrued.

We participate in various managed care sales rebate and other incentive programs, the largest of which relates to Medicaid, Medicare and the Department of Veterans Affairs. Sales rebate and other incentive programs also include contractual volume rebate programs and chargebacks, which are contractual discounts given primarily to federal government agencies, health maintenance organizations, pharmacy benefits managers and group purchasing organizations. We also offer rebate and other incentive programs for our aesthetic products and certain therapeutic products, including *Botox*® Cosmetic, *Juvéderm*®, *Latisse*®, *Acuvail*®, *Aczone*®, *Sanctura XR*® and *Restasis*®, and for certain other skin care products. Sales rebates

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and incentive accruals reduce revenue in the same period that the related sale is recorded and are included in Other accrued expenses in our consolidated balance sheets. The amounts accrued for sales rebates and other incentive programs were $197.0 million and $186.5 million at March 31, 2011 and December 31, 2010, respectively. Provisions for sales rebates and other incentive programs deducted from consolidated sales were $174.5 million and $131.7 million in the first quarter of 2011 and 2010, respectively. The increases in the amounts accrued at March 31, 2011 compared to December 31, 2010 and the provisions for sales rebates and other incentive programs in the first quarter of 2011 compared to the first quarter of 2010 are primarily due to an increase in activity under previously established rebate and incentive programs, principally related to our eye care pharmaceuticals, *Botox*® Cosmetic, urology, skin care and facial aesthetics products, an increase in the number of incentive programs offered, additional contractual discounts to federal government agencies related to the recently enacted health care reform legislation, and increased overall product sales volume. In addition, an increase in our published list prices in the United States for pharmaceutical products, which occurred for several of our products in each of 2011 and 2010, generally results in higher provisions for sales rebates and other incentive programs deducted from consolidated sales.

Our procedures for estimating amounts accrued for sales rebates and other incentive programs at the end of any period are based on available quantitative data and are supplemented by management s judgment with respect to many factors, including but not limited to, current market dynamics, changes in contract terms, changes in sales trends, an evaluation of current laws and regulations and product pricing. Quantitatively, we use historical sales, product utilization and rebate data and apply forecasting techniques in order to estimate our liability amounts. Qualitatively, management s judgment is applied to these items to modify, if appropriate, the estimated liability amounts. There are inherent risks in this process. For example, customers may not achieve assumed utilization levels; customers may misreport their utilization to us; and actual movements of the U.S. Consumer Price Index for All Urban Consumers, or CPI-U, which affect our rebate programs with U.S. federal and state government agencies, may differ from those estimated. On a quarterly basis, adjustments to our estimated liabilities for sales rebates and other incentive programs related to sales made in prior periods have not been material and have generally been less than 0.5% of consolidated product net sales. An adjustment to our estimated liabilities of 0.5% of consolidated product net sales on a quarterly basis would result in an increase or decrease to net sales and earnings before income taxes of approximately $6.0 million to $7.0 million. The sensitivity of our estimates can vary by program and type of customer. Additionally, there is a significant time lag between the date we determine the estimated liability and when we actually pay the liability. Due to this time lag, we record adjustments to our estimated liabilities over several periods, which can result in a net increase to earnings or a net decrease to earnings in those periods. Material differences may result in the amount of revenue we recognize from product sales if the actual amount of rebates and incentives differ materially from the amounts estimated by management.

We recognize license fees, royalties and reimbursement income for services provided as other revenues based on the facts and circumstances of each contractual agreement. In general, we recognize income upon the signing of a contractual agreement that grants rights to products or technology to a third party if we have no further obligation to provide products or services to the third party after entering into the contract. We recognize contingent consideration earned from the achievement of a substantive milestone in its entirety in the period in which the milestone is achieved. We defer income under contractual agreements when we have further obligations that indicate that a separate earnings process has not been completed.

Pensions

We sponsor various pension plans in the United States and abroad in accordance with local laws and regulations. Our U.S. pension plans account for a large majority of our aggregate pension plans net periodic benefit costs and projected benefit obligations. In connection with these plans, we use certain actuarial assumptions to determine the plans net periodic benefit costs and projected benefit obligations, the most significant of which are the expected long-term rate of return on assets and the discount rate.

Our assumption for the weighted average expected long-term rate of return on assets in our U.S. funded pension plan for determining the net periodic benefit cost is 7.25% and 8.25% for 2011 and 2010, respectively. Our assumptions for the weighted average expected long-term rate of return on assets in our non-U.S. funded pension plans are 5.70% and 5.85% for 2011 and 2010, respectively. For our U.S. funded pension plan, we determine, based upon recommendations from our pension plan s investment advisors, the expected rate of return using a building block approach that considers diversification and rebalancing for a long-term portfolio of invested assets. Our investment advisors study historical market returns and preserve long-term historical relationships between equities and fixed income in a manner consistent with the widely-accepted capital market principle that assets with higher volatility generate a greater return over the long run. They also evaluate market factors such as inflation and interest rates before long-term capital market assumptions are determined. For our non-U.S. funded pension plans, the expected rate of return was determined based on asset distribution and assumed long-term rates of return on fixed income instruments and equities. Market conditions and other factors can vary over time and could significantly affect our estimates of the weighted average expected long-term rate of return on plan assets.

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The expected rate of return is applied to the market-related value of plan assets. As a sensitivity measure, the effect of a 0.25% decline in our rate of return on assets assumptions for our U.S. and non-U.S. funded pension plans would increase our expected 2011 pre-tax pension benefit cost by approximately $1.6 million.

The weighted average discount rates used to calculate our U.S. and non-U.S. pension benefit obligations at December 31, 2010 were 5.51% and 5.57%, respectively. The weighted average discount rates used to calculate our U.S. and non-U.S. net periodic benefit costs for 2011 were 5.51% and 5.57%, respectively, and for 2010 were 6.04% and 6.16%, respectively. We determine the discount rate based upon a hypothetical portfolio of high quality fixed income investments with maturities that mirror the pension benefit obligations at the plans measurement date. Market conditions and other factors can vary over time and could significantly affect our estimates for the discount rates used to calculate our pension benefit obligations and net periodic benefit costs for future years. As a sensitivity measure, the effect of a 0.25% decline in the discount rate assumption for our U.S. and non-U.S. pension plans would increase our expected 2011 pre-tax pension benefit costs by approximately $4.1 million and increase our pension plans projected benefit obligations at December 31, 2010 by approximately $34.7 million.

Share-Based Compensation

We recognize compensation expense for all share-based awards made to employees and directors. The fair value of share-based awards is estimated at the grant date using the Black-Scholes option-pricing model and the portion that is ultimately expected to vest is recognized as compensation cost over the requisite service period using the straight-line single option method. The fair value of modifications to share-based awards is generally estimated using a lattice model.

The determination of fair value using the Black-Scholes and lattice option-pricing models is affected by our stock price as well as assumptions regarding a number of complex and subjective variables, including expected stock price volatility, risk-free interest rate, expected dividends and projected employee stock option exercise behaviors. We currently estimate stock price volatility based upon an equal weighting of the historical average over the expected life of the award and the average implied volatility of at-the-money options traded in the open market. We estimate employee stock option exercise behavior based on actual historical exercise activity and assumptions regarding future exercise activity of unexercised, outstanding options.

Share-based compensation expense is recognized only for those awards that are ultimately expected to vest, and we have applied an estimated forfeiture rate to unvested awards for the purpose of calculating compensation cost. These estimates will be revised in future periods if actual forfeitures differ from the estimates. Changes in forfeiture estimates impact compensation cost in the period in which the change in estimate occurs.

Income Taxes

The provision for income taxes is determined using an estimated annual effective tax rate, which is generally less than the U.S. federal statutory rate, primarily because of lower tax rates in certain non-U.S. jurisdictions, research and development, or R&D, tax credits available in the United States, California and other foreign jurisdictions and deductions available in the United States for domestic production activities. Our effective tax rate may be subject to fluctuations during the year as new information is obtained, which may affect the assumptions used to estimate the annual effective tax rate, including factors such as the mix of pre-tax earnings in the various tax jurisdictions in which we operate, valuation allowances against deferred tax assets, the recognition or derecognition of tax benefits related to uncertain tax positions, expected utilization of R&D tax credits and changes in or the interpretation of tax laws in jurisdictions where we conduct business. We recognize deferred tax assets and liabilities for temporary differences between the financial reporting basis and the tax basis of our assets and liabilities along with net operating loss and tax credit carryovers.

We record a valuation allowance against our deferred tax assets to reduce the net carrying value to an amount that we believe is more likely than not to be realized. When we establish or reduce the valuation allowance against our deferred tax assets, our provision for income taxes will increase or decrease, respectively, in the period such determination is made. Valuation allowances against deferred tax assets were $23.8 million and $4.3 million at March 31, 2011 and December 31, 2010, respectively. The increase in the valuation allowance was primarily due to a corresponding increase in a deferred tax asset that we determined required a valuation allowance. Changes in the valuation allowances, when they are recognized in the provision for income taxes, are included as a component of the estimated annual effective tax rate.

We have not provided for withholding and U.S. taxes for the unremitted earnings of certain non-U.S. subsidiaries because we have currently reinvested these earnings indefinitely in these foreign operations. At December 31, 2010, we had approximately $2,109.4 million in unremitted earnings outside the United States for which withholding and U.S. taxes were not provided. Income tax expense would be incurred if these funds were remitted to the United States. It is not practicable to estimate the amount of the deferred tax liability on such unremitted earnings. Upon remittance, certain foreign countries

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impose withholding taxes that are then available, subject to certain limitations, for use as credits against our U.S. tax liability, if any. We annually update our estimate of unremitted earnings outside the United States after the completion of each fiscal year.

Acquisitions

The accounting for acquisitions requires extensive use of estimates and judgments to measure the fair value of the identifiable tangible and intangible assets acquired, including in-process research and development, and liabilities assumed. Additionally, we must determine whether an acquired entity is considered to be a business or a set of net assets, because the excess of the purchase price over the fair value of net assets acquired can only be recognized as goodwill in a business combination.

On July 1, 2010, we completed a business combination agreement and effected a revised distribution agreement with our distributor in Turkey. We paid $33.0 million for the termination of the original distribution agreement and purchased the commercial assets related to the selling of our products in Turkey for $6.1 million in cash and estimated contingent consideration of $36.7 million as of the acquisition date. On January 15, 2010, we acquired Serica Technologies, Inc., or Serica, for an aggregate purchase price of approximately $63.7 million, net of cash acquired. We accounted for these acquisitions as business combinations. The tangible and intangible assets acquired and liabilities assumed in connection with these acquisitions were recognized based on their estimated fair values at the acquisition dates. The determination of estimated fair values requires significant estimates and assumptions including, but not limited to, determining the timing and estimated costs to complete the in-process projects, projecting regulatory approvals, estimating future cash flows and developing appropriate discount rates. We believe the estimated fair values assigned to the assets acquired and liabilities assumed are based on reasonable assumptions.

Impairment Evaluations for Goodwill and Purchased Intangible Assets

We evaluate goodwill for impairment on an annual basis, or more frequently if we believe indicators of impairment exist, by comparing the carrying value of each of our reporting units to their estimated fair value. We have identified two reporting units, specialty pharmaceuticals and medical devices, and currently perform our annual evaluation as of October 1 each year.

We primarily use the income approach and the market approach to valuation that include the discounted cash flow method, the guideline company method, as well as other generally accepted valuation methodologies to determine the fair value of our reporting units. Upon completion of the October 2010 annual impairment assessment, we determined that no impairment was indicated as the estimated fair value of each of the two reporting units exceeded its respective carrying value. As of March 31, 2011, we do not believe any significant indicators of impairment exist for our goodwill that would require additional analysis.

We also review purchased intangible assets for impairment when events or changes in circumstances indicate that the carrying value of our intangible assets may not be recoverable. An impairment in the carrying value of an intangible asset is recognized whenever anticipated future undiscounted cash flows from an intangible asset are estimated to be less than its carrying value.

In March 2011, we decided to discontinue development of the *EasyBand* Remote Adjustable Gastric Band System, or *EasyBand*, a technology that we acquired in connection with our 2007 acquisition of EndoArt SA, or EndoArt. As a result, in the first quarter of 2011 we recorded a pre-tax impairment charge of $16.1 million for the intangible assets associated with the *EasyBand* technology.

Significant management judgment is required in the forecasts of future operating results that are used in our impairment evaluations. The estimates we have used are consistent with the plans and estimates that we use to manage our business. It is possible, however, that the plans may change and estimates used may prove to be inaccurate. If our actual results, or the plans and estimates used in future impairment analyses, are lower than the original estimates used to assess the recoverability of these assets, we could incur future impairment charges.

Operations

Headquartered in Irvine, California, we are a multi-specialty health care company focused on discovering, developing and commercializing innovative pharmaceuticals, biologics, medical devices and over-the-counter products that enable people to live life to its greatest potential to see more clearly, move more freely and express themselves more fully. Our diversified approach enables us to follow our research and development into new specialty areas where unmet needs are significant.

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We discover, develop and commercialize specialty pharmaceutical, biologics, medical devices and over-the-counter products for the ophthalmic, neurological, medical aesthetics, medical dermatology, breast aesthetics, obesity intervention, urological and other specialty markets in more than 100 countries around the world. We are a pioneer in specialty pharmaceutical research, targeting products and technologies related to specific disease areas such as chronic dry eye, glaucoma, retinal disease, psoriasis, acne, movement disorders, neuropathic pain and genitourinary diseases. Additionally, we are a leader in discovering, developing and marketing therapeutic and aesthetic biological, pharmaceutical and medical device products, including saline and silicone gel breast implants, dermal fillers and obesity intervention products. At March 31, 2011, we employed approximately 9,500 persons around the world. Our principal markets are the United States, Europe, Latin America and Asia Pacific.

Results of Operations

We operate our business on the basis of two reportable segments specialty pharmaceuticals and medical devices. The specialty pharmaceuticals segment produces a broad range of pharmaceutical products, including: ophthalmic products for dry eye, glaucoma, retinal diseases and ocular surface disease; *Botox*® for certain therapeutic and aesthetic indications; skin care products for acne, psoriasis, eyelash growth and other prescription and over-the-counter skin care products; and urologics products. The medical devices segment produces a broad range of medical devices, including: breast implants for augmentation, revision and reconstructive surgery; obesity intervention products, including the *Lap-Band*® System and the *Orbera* Intragastric Balloon System; and facial aesthetics products. We provide global marketing strategy teams to coordinate the development and execution of a consistent marketing strategy for our products in all geographic regions that share similar distribution channels and customers.

Management evaluates our business segments and various global product portfolios on a revenue basis, which is presented below in accordance with GAAP. We also report sales performance using the non-GAAP financial measure of constant currency sales. Constant currency sales represent current period reported sales, adjusted for the translation effect of changes in average foreign exchange rates between the current period and the corresponding period in the prior year. We calculate the currency effect by comparing adjusted current period reported sales, calculated using the monthly average foreign exchange rates for the corresponding period in the prior year, to the actual current period reported sales. We routinely evaluate our net sales performance at constant currency so that sales results can be viewed without the impact of changing foreign currency exchange rates, thereby facilitating period-to-period comparisons of our sales. Generally, when the U.S. dollar either strengthens or weakens against other currencies, the growth at constant currency rates will be higher or lower, respectively, than growth reported at actual exchange rates.

The following table compares net sales by product line within each reportable segment and certain selected pharmaceutical products for the three month periods ended March 31, 2011 and 2010:

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	Three months ended March 31, 2011	March 31, 2010	Change in Product Net Sales Total (in millions)	Performance	Currency	Percent Change in Product Net Sales Total	Performance	Currency
Net Sales by Product Line:								
Specialty Pharmaceuticals:								
Eye Care Pharmaceuticals	$ 591.9	$ 512.0	$ 79.9	$ 75.2	$ 4.7	15.6 %	14.7 %	0.9%
Botox®/Neuromodulator	364.5	331.0	33.5	28.9	4.6	10.1 %	8.7 %	1.4%
Skin Care	58.7	50.6	8.1	8.0	0.1	16.0 %	15.8 %	0.2%
Urologics	13.3	13.7	(0.4)	(0.4)		(2.9)%	(2.9)%	%
Total Specialty Pharmaceuticals	1,028.4	907.3	121.1	111.7	9.4	13.3 %	12.3 %	1.0%
Medical Devices:								
Breast Aesthetics	84.1	77.9	6.2	5.5	0.7	8.0 %	7.1 %	0.9%
Obesity Intervention	52.1	61.2	(9.1)	(9.8)	0.7	(14.9)%	(16.0)%	1.1%
Facial Aesthetics	88.2	59.4	28.8	27.9	0.9	48.5 %	47.0 %	1.5%
Total Medical Devices	224.4	198.5	25.9	23.6	2.3	13.0 %	11.9 %	1.1%
Total product net sales	$ 1,252.8	$ 1,105.8	$ 147.0	$ 135.3	$ 11.7	13.3 %	12.2 %	1.1%
Domestic product net sales	60.9%	62.5%						
International product net sales	39.1%	37.5%						
Selected Product Net Sales (a):								
Alphagan® *P*, *Alphagan*® and *Combigan*®	$ 100.2	$ 94.1	$ 6.1	$ 5.4	$ 0.7	6.5 %	5.8 %	0.7%
Lumigan® Franchise	142.2	119.6	22.6	21.8	0.8	18.9 %	18.2 %	0.7%
Restasis®	161.4	133.4	28.0	27.9	0.1	21.0 %	20.9 %	0.1%
Sanctura® Franchise	13.3	13.7	(0.4)	(0.4)		(2.9)%	(2.9)%	%
Latisse®	25.3	18.8	6.5	6.4	0.1	34.2 %	33.6 %	0.6%

(a) Percentage change in selected product net sales is calculated on amounts reported to the nearest whole dollar.

Product Net Sales

Product net sales increased by $147.0 million in the first quarter of 2011 compared to the first quarter of 2010 due to an increase of $121.1 million in our specialty pharmaceuticals product net sales and an increase of $25.9 million in our medical devices product net sales. The increase in specialty pharmaceuticals product net sales is due to increases in product net sales of our eye care pharmaceuticals, *Botox*®, and skin care product lines, partially offset by a small decrease in product net sales of our urologics product line. The increase in medical devices product net sales reflects an increase in product net sales of our breast aesthetics and facial aesthetics product lines, partially offset by a decrease in product net sales of our obesity intervention product line.

Several of our products, including *Botox*® Cosmetic, *Latisse*® and our facial aesthetics, obesity intervention and breast implant products, are purchased based on consumer choice and have limited reimbursement or are not reimbursable by government or other health care plans and are, therefore, partially or wholly paid for directly by the consumer. As such, the general economic environment and level of consumer spending have a significant effect on our sales of these products.

In March 2010, the U.S. government enacted the Patient Protection and Affordable Care Act, as amended by the Health Care and Education Affordability Reconciliation Act, collectively, the PPACA, reforming the U.S. health care system. The PPACA includes provisions that have a significant negative impact on our product net sales, including an extension of Medicaid and Medicare benefits to new patient populations, an increase in Medicaid rebates owed by manufacturers under the Medicaid Drug Rebate Program and a future increase in the initial coverage limit for Medicare participants. In the first quarter of 2011, the additional rebates related to the PPACA had a negative impact of approximately $10.0 million on our product net sales. The PPACA did not impact product net sales in the first quarter of 2010. Based on internal information and

assumptions, we currently estimate that the PPACA will have a negative impact on our fiscal year 2011 product net sales of approximately $40.0 million. The PPACA also established an annual non-deductible fee on entities that sell branded prescription drugs or biologics to specified government programs in the United States. We expect this fee will have a negative impact on our selling, general and administrative expenses of approximately $20.4 million in 2011. In addition, we expect incremental price reductions and rebate increases mandated by European governments to have a negative impact on our 2011 product net sales of approximately $30.0 million. In the aggregate, we expect that incremental costs of healthcare

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reform under the PPACA and the effect of European pricing pressures will have a negative impact on our fiscal year 2011 earnings on a pre-tax equivalent basis of approximately $100.0 million.

Eye care pharmaceuticals product net sales increased in the first quarter of 2011 compared to the first quarter of 2010 primarily due to an increase in net sales of *Restasis*®, our therapeutic treatment for chronic dry eye disease, an increase in new product sales of our glaucoma drug *Lumigan*® 0.01%, which was launched in the United States in the fourth quarter of 2010, an increase in sales of *Ganfort*, our *Lumigan*® and timolol combination for the treatment of glaucoma, an increase in sales of *Combigan*®, our *Alphagan*® and timolol combination for the treatment of glaucoma, an increase in sales of our glaucoma drug *Alphagan*® *P* 0.1%, an increase in sales of *Ozurdex*®, our biodegradable, sustained-release steroid implant for the treatment of certain retinal diseases, an increase in new product sales of *Zymaxid*®, our next-generation anti-infective product in the fluoroquinolone category indicated for the treatment of bacterial conjunctivitis, which was launched in the second quarter of 2010, an increase in new product sales of *Lastacaft*, our topical allergy medication for the treatment and prevention of itching associated with allergic conjunctivitis, which we launched in the United States in January 2011, and an increase in sales of our artificial tears products *Refresh*® and *Refresh*® *Optive*, partially offset by decreases in sales of our glaucoma drugs *Alphagan*®*, Alphagan*® *P* 0.15% and *Lumigan*® 0.03%, our older-generation fluoroquinolone *Zymar*® and our non-steroidal anti-inflammatory drug *Acuvail*®. Beginning in February 2011 we discontinued the U.S. distribution of *Zymar*®.

In May 2011 a generic version of *Elestat*®, our older generation topical antihistamine used for the prevention of itching associated with allergic conjunctivitis, was launched in the United States. In addition, we expect a generic version of *Zymar*® to be launched in the United States during 2011. While we estimate that our product net sales will be negatively impacted in 2011 due to sales of generic formulations of these products, we expect that any such negative impact on product net sales will be partially offset by increased sales of *Lastacaft* and *Zymaxid*®.

We increased prices on certain eye care pharmaceutical products in the United States in the second half of 2010 and the first quarter of 2011. Effective January 8, 2011, we increased the published U.S. list price for *Restasis*®, *Alphagan*® *P* 0.1%, *Alphagan*® *P* 0.15%, *Combigan*®*, Zymar*®*, Zymaxid*®*, Acular*®*, Acular LS*® and *Acuvail*® by four percent and *Lumigan*® 0.1% and *Lumigan*® 0.3% by eight percent. These price increases had a positive net effect on our U.S. sales in the first quarter of 2011 compared to the first quarter of 2010, but the actual net effect is difficult to determine due to the various managed care sales rebate and other incentive programs in which we participate. Wholesaler buying patterns and the change in dollar value of the prescription product mix also affected our reported net sales dollars, although we are unable to determine the impact of these effects.

Total sales of *Botox*® increased in the first quarter of 2011 compared to the first quarter of 2010 due to strong growth in sales for both cosmetic and therapeutic use in the United States, Canada, Latin America, Asia Pacific and, to a lesser degree, Europe. We believe our worldwide market share for neuromodulators, including *Botox*®, was approximately 80% in the fourth quarter of 2010, the last quarter for which market data is available.

Skin care product net sales increased in the first quarter of 2011 compared to the first quarter of 2010 primarily due to an increase in sales of *Latisse*®, our treatment for inadequate or insufficient eyelashes, and an increase in sales of *Aczone*®, our topical dapsone treatment for acne vulgaris, partially offset by a decrease in total sales of *Tazorac*®, *Zorac*® and *Avage*®, our topical tazarotene products. Effective January 8, 2011, we increased the published U.S. list price for *Aczone*® by approximately four percent, and *Tazorac*® and *Avage*® by approximately fifteen percent. We expect a generic version of our *Tazorac*® cream product to be launched in the United States in mid-2011 and estimate that our product net sales will be negatively impacted in 2011 due to sales of generic formulations of this product.

Urologics sales, which are presently concentrated in the United States and consist of our *Sanctura*® franchise products for the treatment of overactive bladder, decreased in the first quarter of 2011 compared to the first quarter of 2010, primarily due to lower sales of *Sanctura*®, our twice-a-day anticholinergic for the treatment of overactive bladder, or OAB, which was negatively impacted by the launch of trospium chloride generics in September 2010, partially offset by an increase in sales of *Sanctura XR*®, our second generation, once-daily anticholinergic for the treatment of OAB. Effective January 8, 2011, we increased the published U.S. list price for *Sanctura XR*® by eight percent and *Sanctura*® by ten percent.

We have a policy to attempt to maintain average U.S. wholesaler inventory levels of our specialty pharmaceutical products at an amount less than eight weeks of our net sales. At March 31, 2011, based on available external and internal information, we believe the amount of average U.S. wholesaler inventories of our specialty pharmaceutical products was near the lower end of our stated policy levels.

Breast aesthetics product net sales, which consist primarily of sales of silicone gel and saline breast implants and tissue expanders, increased in the first quarter of 2011 compared to the first quarter of 2010 due to increases in sales in all of our principal geographic markets. The increase in sales of breast aesthetics products was primarily due to higher silicone gel

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implant and tissue expander unit volume, and the continued transition of the U.S. market to higher priced silicone gel products from lower priced saline products.

Obesity intervention product net sales, which consist primarily of sales of devices used for minimally invasive long-term treatments of obesity such as our *Lap-Band*® and *Lap-Band AP*® Systems and *Orbera* System, decreased in the first quarter of 2011 compared to the first quarter of 2010 primarily due to a decrease in sales in the United States and Australia, and a small decrease in sales in Europe, partially offset by an increase in sales in Latin America. We believe sales of obesity intervention products in the United States and other principal geographic markets continued to be negatively impacted by general economic conditions given the substantial patient co-pays associated with these products and government spending restrictions.

Facial aesthetics product net sales, which consist primarily of sales of hyaluronic acid-based dermal fillers used to correct facial wrinkles, increased in the first quarter of 2011 compared to the first quarter of 2010 primarily due to significant increases in sales in the United States and all of our other principal geographic markets. We believe the increase in sales of facial aesthetic products was primarily due to the February 2010 launch of *Juvéderm*® XC with lidocaine in the United States and recent launches of *Juvéderm*® with lidocaine and *Juvéderm*® *Voluma* in other international markets and a global expansion of the dermal filler market, partially offset by a decline in sales of older generation collagen-based dermal fillers, which we discontinued selling in early 2011.

Foreign currency changes increased product net sales by $11.7 million in the first quarter of 2011 compared to the first quarter of 2010, primarily due to the strengthening of the Canadian dollar, Brazilian real, Australian dollar and U.K. pound compared to the U.S. dollar, partially offset by a slight weakening of the euro compared to the U.S. dollar.

U.S. product net sales as a percentage of total product net sales decreased by 1.6 percentage points to 60.9% in the first quarter of 2011 compared to U.S. sales of 62.5% in the first quarter of 2010, due primarily to higher sales growth in our international markets compared to the U.S. market for our eye care pharmaceuticals and *Botox*® product lines and a higher percentage decline in sales in the U.S. market compared to our total international markets for our obesity intervention product line, partially offset by an increase in sales of our skin care products, which are highly concentrated in the United States. Additionally, international sales benefited from a positive translation impact due to a general strengthening of foreign currencies compared to the U.S. dollar in markets where we sold products in the first quarter of 2011 compared to the first quarter of 2010.

Other Revenues

Other revenues decreased $30.5 million to $18.4 million in the first quarter of 2011 compared to $48.9 million in the first quarter of 2010. The decrease in other revenues is primarily related to the prior year impact of an upfront net licensing fee of $36.0 million that we recognized in the first quarter of 2010 related to an agreement with Bristol-Myers Squibb Company for the exclusive worldwide rights to develop, manufacture and commercialize an investigational medicine for neuropathic pain, partially offset by an increase in royalty income in the first quarter of 2011 compared to the first quarter of 2010 from sales of brimonidine products by Alcon, Inc. in the United States under a licensing agreement and an increase in royalty income from sales of *Lumigan*® by Senju Pharmaceutical Co., Ltd., or Senju, in Japan under a licensing agreement.

Cost of Sales

Cost of sales increased $13.1 million, or 7.7%, in the first quarter of 2011 to $183.3 million, or 14.6% of product net sales, compared to $170.2 million, or 15.4% of product net sales in the first quarter of 2010. This increase in cost of sales primarily resulted from the 13.3% increase in total product net sales and an increase in cost of sales as a percentage of product net sales for our facial aesthetics products due to an increase in provisions for inventory returns, partially offset by a decrease in cost of sales as a percentage of product net sales for our eye care pharmaceuticals, skin care, *Botox*® and breast aesthetics product lines primarily due to volume-based manufacturing efficiencies and lower royalty expenses.

Selling, General and Administrative

Selling, general and administrative, or SG&A, expenses increased $115.7 million, or 24.4%, to $589.5 million, or 47.1% of product net sales, in the first quarter of 2011 compared to $473.8 million, or 42.8% of product net sales, in the first quarter of 2010. SG&A expenses in the first quarter of 2011 include an upfront payment of $60.0 million related to a collaboration and co-promotion agreement with MAP Pharmaceuticals, Inc., or MAP, for the development and commercialization of *Levadex*, a self-administered, orally inhaled therapy for the treatment of acute migraine in adults that has not yet achieved regulatory approval and other potential indications in the United States, a gain of $9.4 million from the substantially complete liquidation of a foreign subsidiary and fixed asset impairment charges of $2.3 million related to the discontinued development of *EasyBand*, and $1.6 million of stockholder derivative litigation costs associated with the 2010

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global settlement with the U.S. Department of Justice, or DOJ, regarding our past U.S. sales and marketing practices relating to certain therapeutic uses of *Botox*®. SG&A expenses in the first quarter of 2010 include $4.5 million of costs associated with the DOJ investigation that related to sales and marketing practices in connection with *Botox*®. Excluding the effect of the items described above, SG&A expenses increased $65.7 million, or 14.0%, to $535.0 million, or 42.7% of product net sales, in the first quarter of 2011 compared to $469.3 million, or 42.4% of product net sales in the first quarter of 2010. The increase in SG&A expenses in dollars, excluding the charges described above, primarily relates to increases in selling, marketing, promotion and general and administrative expenses and the negative translation impact due to a general strengthening of foreign currencies compared to the U.S. dollar. The increase in selling and marketing expenses in the first quarter of 2011 compared to the first quarter of 2010 principally relates to increased personnel and related incentive compensation costs that support the 13.3% increase in product net sales, and additional costs supporting the expansion of our sales forces, including the addition of new direct operations in Turkey, Poland and the Philippines. The increase in promotion expenses is primarily due to increased direct-to-consumer advertising for *Restasis*® and *Juvéderm*®, partially offset by a decline in direct-to-consumer advertising for *Latisse*®. The increase in general and administrative expenses is primarily due to the negative impact of the fee established by the PPACA for selling branded pharmaceuticals to certain U.S. government programs, increased compliance costs associated with the Corporate Integrity Agreement entered into in 2010 with the Office of Inspector General of the Department of Health and Human Services, an increase in losses from the disposal of fixed assets, and an increase in incentive compensation costs, information systems, finance and human resource administrative costs. The increase in SG&A expenses as a percentage of product net sales, excluding the items described above, in the first quarter of 2011 compared to the first quarter of 2010 is primarily due to the higher 14.0% increase in SG&A expenses relative to the 13.3% increase in product net sales during the same period.

Research and Development

Research and development, or R&D, expenses decreased $25.0 million, or 11.2%, to $197.7 million in the first quarter of 2011, or 15.8% of product net sales, compared to $222.7 million, or 20.1% of product net sales in the first quarter of 2010. R&D expenses in the first quarter of 2010 included a charge of $43.0 million for an upfront payment for the in-licensing of technology for treatment of nocturia, a urological disorder characterized by frequent urination at nighttime, from Serenity Pharmaceuticals, LLC, or Serenity, that has not yet achieved regulatory approval. Excluding the effect of this charge, R&D expenses increased by $18.0 million, or 10.0% in the first quarter of 2011 compared to the first quarter of 2010. The increase in R&D expenses, excluding the 2010 upfront payment charge to Serenity, was primarily due to increased spending on new technology discovery programs, next generation eye care pharmaceuticals products for the treatment of glaucoma and retinal diseases, potential new treatment applications for *Latisse*®, *Botox*® for the treatment of overactive bladder, hyaluronic-acid based dermal filler products, tissue regeneration technology acquired in the Serica acquisition and obesity intervention products, partially offset by a reduction in expenses related to the development of *Ozurdex*® and a small decrease in spending for certain urology products.

Amortization of Acquired Intangible Assets

Amortization of acquired intangible assets decreased $4.6 million to $32.5 million in the first quarter of 2011, or 2.6% of product net sales, compared to $37.1 million, or 3.4% of product net sales, in the first quarter of 2010. The decrease in amortization expense in dollars and as a percentage of product net sales is primarily due to the impairment of the *Sanctura*® intangible assets in the third quarter of 2010, partially offset by an increase in the balance of intangible assets subject to amortization, including a capitalized upfront licensing payment in September 2010 for *Lastacaft*, an eye care product previously approved for marketing, licensing assets related to *Botox*® Cosmetic distribution rights in Japan and China that we reacquired from GlaxoSmithKline in the first quarter of 2010 and other intangible assets that we acquired in connection with our July 2010 purchase of our distributor s business related to our products in Turkey.

Intangible Asset Impairment

In March 2011, we decided to discontinue development of *EasyBand*, a technology that we acquired in connection with our 2007 acquisition of EndoArt. As a result, in the first quarter of 2011 we recorded a pre-tax impairment charge of $16.1 million for the intangible assets associated with the *EasyBand* technology.

Restructuring Charges

Restructuring charges were $4.6 million and $0.6 million in the first quarter of 2011 and 2010, respectively.

Discontinued Development of *EasyBand*

In March 2011, we decided to discontinue development of the *EasyBand* Remote Adjustable Gastric Band System and close the related research and development facility in Switzerland.

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As a result of discontinuing the development of *EasyBand* and the closure of the related research and development facility, in the first quarter of 2011 we recorded a pre-tax impairment charge of $16.1 million for the intangible assets associated with the *EasyBand* technology, fixed asset impairment charges of $2.3 million and a gain of $9.4 million from the substantially complete liquidation of our investment in a foreign subsidiary. In addition, we recorded $4.6 million of restructuring charges, consisting of $3.0 million of employee severance and other one-time termination benefits for approximately 30 people affected by the facility closure, $1.5 million of contract termination costs and $0.1 million of other related costs.

Other Restructuring Activities and Integration Costs

We did not incur any other restructuring charges during the three month period ended March 31, 2011.

Included in the three month period ended March 31, 2010 are $0.1 million of restructuring charges primarily for employee severance and other one-time termination benefits related to our fiscal year 2009 restructuring plan, $0.4 million of restructuring charges primarily for employee severance related to our acquisition of Serica and $0.1 million of restructuring charges for an abandoned leased facility related to our fiscal year 2005 restructuring and streamlining of our European operations.

Included in the three month period ended March 31, 2011 are $0.6 million of SG&A expenses related to transaction costs associated with the collaboration and co-promotion agreements with MAP, $0.2 million of SG&A expenses related to integration and transaction costs associated with the purchase of our distributor s business related to our products in Turkey and $0.2 million of SG&A expenses related to transaction costs associated with the announced purchase of our distributor s business related to our products in South Africa.

Included in the three month period ended March 31, 2010 are $0.5 million of SG&A expenses related to integration and transaction costs associated with our acquisition of Serica, $0.2 million of SG&A expenses related to transaction costs associated with the purchase of our distributor s business related to our products in Turkey and $0.3 million of SG&A expenses related to transaction costs associated with the license, development and commercialization agreement with Serenity.

Operating Income

Management evaluates business segment performance on an operating income basis exclusive of general and administrative expenses and other indirect costs, legal settlement expenses, intangible asset impairment and related costs, restructuring charges, in-process research and development expenses, amortization of certain identifiable intangible assets related to business combinations and asset acquisitions and related capitalized licensing costs and certain other adjustments, which are not allocated to our business segments for performance assessment by our chief operating decision maker. Other adjustments excluded from our business segments for purposes of performance assessment represent income or expenses that do not reflect, according to established Company-defined criteria, operating income or expenses associated with our core business activities.

For the first quarter of 2011, general and administrative expenses, other indirect costs and other adjustments not allocated to our business segments for purposes of performance assessment consisted of general and administrative expenses of $90.6 million, an upfront payment of $60.0 million to MAP for a collaboration and co-promotion agreement related to technology that has not achieved regulatory approval and related transaction costs of $0.6 million, fixed asset impairment charges of $2.3 million, a gain of $9.4 million from the substantially complete liquidation of the Company s investment in a foreign subsidiary, stockholder derivative litigation costs of $1.6 million in connection with the global settlement with the DOJ regarding our past U.S. sales and marketing practices relating to *Botox*®, integration and transaction costs of $0.4 million associated with the purchases of our distributors businesses related to our products in Turkey and South Africa, and other net indirect costs of $10.8 million.

For the first quarter of 2010, general and administrative expenses, other indirect costs and other adjustments not allocated to our business segments for purposes of performance assessment consisted of licensing fee income of $36.0 million for a development and commercialization agreement with Bristol-Myers Squibb Company, general and administrative expenses of $82.5 million, costs associated with the DOJ investigation regarding our past U.S. sales and marketing practices relating to *Botox*® of $4.5 million, an upfront licensing fee included in R&D expenses of $43.0 million paid to Serenity for technology that has not achieved regulatory approval and related transaction costs of $0.3 million, integration and transaction costs of $0.5 million related to our acquisition of Serica, transaction costs of $0.2 million associated with the purchase of our distributor s business related to our products in Turkey, and other net indirect costs of $1.7 million.

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The following table presents operating income for each reportable segment for the three month periods ended March 31, 2011 and 2010 and a reconciliation of our segments operating income to consolidated operating income:

	Three months ended	
	March 31, 2011	March 31, 2010
	(in millions)	
Operating income:		
Specialty pharmaceuticals	$ 384.2	$ 311.9
Medical devices	67.5	67.1
Total segments	451.7	379.0
General and administrative expenses, other indirect costs and other adjustments	156.9	96.7
Amortization of acquired intangible assets (a)	26.6	31.4
Intangible asset impairment	16.1	
Restructuring charges	4.6	0.6
Total operating income	$ 247.5	$ 250.3

(a) Represents amortization of certain identifiable intangible assets related to business combinations and asset acquisitions and related capitalized licensing costs, as applicable.

Our consolidated operating income in the first quarter of 2011 was $247.5 million, or 19.8% of product net sales, compared to consolidated operating income of $250.3 million, or 22.6% of product net sales in the first quarter of 2010. The $2.8 million decrease in consolidated operating income was due to a $16.1 million intangible asset impairment charge, a $30.5 million decrease in other revenues, a $13.1 million increase in cost of sales, a $115.7 million increase in SG&A expenses and a $4.0 million increase in restructuring charges, partially offset by a $147.0 million increase in product net sales, a $25.0 million decrease in R&D expenses and a $4.6 million decrease in amortization of acquired intangible assets.

Our specialty pharmaceuticals segment operating income in the first quarter of 2011 was $384.2 million, compared to operating income of $311.9 million in the first quarter of 2010. The $72.3 million increase in our specialty pharmaceuticals segment operating income was due primarily to an increase in product net sales of our eye care pharmaceuticals, *Botox*® and skin care product lines, partially offset by an increase in selling and marketing expenses and an increase in R&D expenses.

Our medical devices segment operating income in the first quarter of 2011 was $67.5 million, compared to operating income of $67.1 million in the first quarter of 2010. The $0.4 million increase in our medical devices segment operating income was due primarily to an increase in product net sales of our breast aesthetics and facial aesthetics product lines, partially offset by a decrease in product net sales of our obesity intervention product line, an increase in overall promotion and selling expenses and an increase in R&D expenses.

Non-Operating Income and Expense

Total net non-operating expense in the first quarter of 2011 was $32.3 million compared to $18.3 million in the first quarter of 2010. Interest income in the first quarter of 2011 was $2.3 million compared to interest income of $1.3 million in the first quarter of 2010. The increase in interest income was primarily due to higher average cash equivalent and short-term investment balances earning interest of approximately $778 million in the first quarter of 2011 compared to the first quarter of 2010. Interest expense increased $8.1 million to $24.7 million in the first quarter of 2011 compared to $16.6 million in the first quarter of 2010. Interest expense increased primarily due to the issuance in September 2010 of our 3.375% Senior Notes due 2020, or 2020 Notes, and a charge for statutory interest expense in the first quarter of 2011, compared to a reversal of previously accrued statutory interest expense resulting from a change in estimate related to uncertain tax positions in the first quarter of 2010. Other, net expense was $9.9 million in the first quarter of 2011, consisting primarily of a net unrealized loss on derivative instruments of $6.9 million and $4.0 million in net realized losses from foreign currency transactions, partially offset by a gain of $0.5 million on the sale of a third party equity investment. Other, net expense was $3.0 million in the first quarter of 2010, consisting primarily of a net unrealized loss on derivative instruments of $0.7 million and $2.4 million in net realized losses from foreign currency transactions.

Income Taxes

Our effective tax rate for the first quarter of 2011 was 26.2%. Included in our earnings before income taxes for the first quarter of 2011 are a $60.0 million upfront payment related to a collaboration and co-promotion agreement with MAP, an intangible asset impairment charge of $16.1 million, restructuring charges of $4.6 million, fixed asset impairment charges of $2.3 million and a gain of $9.4 million from the substantially complete liquidation of a foreign subsidiary resulting from the

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discontinued development of *EasyBand*. In the first quarter of 2011, we recorded income tax benefits of $22.2 million associated with the upfront payment related to the collaboration and co-promotion agreement with MAP. In the first quarter of 2011, we did not record any tax benefits related to the intangible asset impairment charge, restructuring charges, fixed asset impairment charges and the gain from the substantially complete liquidation of our investment in a foreign subsidiary resulting from the discontinued development of *EasyBand* since we do not expect to be able to utilize tax deductions in the jurisdiction where these costs were incurred. Excluding the impact of the net pre-tax charges of $73.6 million and the net income tax benefits of $22.2 million for the items discussed above, our adjusted effective tax rate for the first quarter of 2011 was 27.2%. We believe that the use of an adjusted effective tax rate provides a more meaningful measure of the impact of income taxes on our results of operations because it excludes the effect of certain items that are not included as part of our core business activities. This allows investors to better determine the effective tax rate associated with our core business activities.

The calculation of our adjusted effective tax rate for the first quarter of 2011 is summarized below:

	(in millions)
Earnings before income taxes, as reported	$ 215.2
Upfront payment for a collaboration and co-promotion agreement with MAP	60.0
Restructuring charges	4.6
Aggregate net expense for the fixed asset impairment, gain from the substantially complete liquidation of a foreign subsidiary and intangible asset impairment resulting from the discontinued development of *Easyband*	9.0
	$ 288.8
Provision for income taxes, as reported	$ 56.4
Income tax benefit for:	
Upfront payment for a collaboration and co-promotion agreement with MAP	22.2
	$ 78.6
Adjusted effective tax rate	27.2%

Our effective tax rate for the first quarter of 2010 was 27.2%. Our effective tax rate for the year ended December 31, 2010 was 97.1% and our adjusted effective tax rate for the year ended December 31, 2010 was 28.0%. Included in our earnings before income taxes for 2010 are total pre-tax charges of $609.2 million in connection with the global settlement with the DOJ regarding our past U.S. sales and marketing practices relating to certain therapeutic uses of *Botox*®, a $369.1 million aggregate charge related to the impairment of the *Sanctura*® Assets and related costs, a $33.0 million charge related to the termination of a distributor agreement in Turkey, a $43.0 million charge for an upfront payment for technology that has not achieved regulatory approval, restructuring charges of $0.3 million and license fee income of $36.0 million related to an upfront fee for product rights we licensed to Bristol-Myers Squibb Company. In 2010, we recorded income tax benefits of $21.4 million related to the global settlement with the DOJ regarding our past U.S. sales and marketing practices relating to certain therapeutic uses of *Botox*®, $140.5 million related to the impairment of the *Sanctura*® Assets and related costs, $2.8 million related to the termination of a distributor agreement in Turkey, $15.6 million related to the upfront payment for technology that has not achieved regulatory approval and $0.2 million related to the restructuring charges, and an income tax expense of $13.7 million related to the upfront license fee income. Excluding the impact of the net pre-tax charges of $1,018.6 million and the net income tax benefits of $166.8 million for the items discussed above, our adjusted effective tax rate for 2010 was 28.0%.

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The calculation of our adjusted effective tax rate for the year ended December 31, 2010 is summarized below:

	(in millions)
Earnings before income taxes, as reported	$ 170.8
Settlement with the DOJ related to U.S. sales and marketing practices for *Botox*®	609.2
Impairment of the *Sanctura*® Assets and related costs	369.1
Termination of a distributor agreement in Turkey	33.0
Upfront payment for technology that has not achieved regulatory approval	43.0
Restructuring charges	0.3
Upfront license fee income	(36.0)
	$ 1,189.4
Provision for income taxes, as reported	$ 165.9
Income tax benefit (provision) for:	
Settlement with the DOJ related to U.S. sales and marketing practices for *Botox*®	21.4
Impairment of the *Sanctura*® Assets and related costs	140.5
Termination of a distributor agreement in Turkey	2.8
Upfront payment for technology that has not achieved regulatory approval	15.6
Restructuring charges	0.2
Upfront license fee income	(13.7)
	$ 332.7
Adjusted effective tax rate	28.0%

The decrease in the adjusted effective tax rate to 27.2% in the first quarter of 2011 compared to the adjusted effective tax rate for the year ended December 31, 2010 of 28.0% is primarily due to the detrimental tax rate effect in 2010 of changes in our deferred tax asset and liability balances related to a change in California tax law and an increase in 2011 compared to 2010 in the expected mix of annual earnings in international jurisdictions, which generally have lower tax rates than the United States.

Net Earnings Attributable to Noncontrolling Interest

Our net earnings attributable to noncontrolling interest for our majority-owned subsidiaries were $0.5 million and $1.1 million in the first quarter of 2011 and 2010, respectively.

Net Earnings Attributable to Allergan, Inc.

Our net earnings attributable to Allergan, Inc. in the first quarter of 2011 was $158.3 million compared to net earnings attributable to Allergan, Inc. of $167.9 million in the first quarter of 2010. The $9.6 million decrease in net earnings attributable to Allergan, Inc. was primarily the result of the decrease in operating income of $2.8 million and the increase in net non-operating expense of $14.0 million, partially offset by the decrease in the provision for income taxes of $6.6 million and the decrease in net earnings attributable to noncontrolling interest of $0.6 million.

Liquidity and Capital Resources

We assess our liquidity by our ability to generate cash to fund our operations. Significant factors in the management of liquidity are: funds generated by operations; levels of accounts receivable, inventories, accounts payable and capital expenditures; the extent of our stock repurchase program; funds required for acquisitions and other transactions; funds available under our credit facilities; and financial flexibility to attract long-term capital on satisfactory terms.

Historically, we have generated cash from operations in excess of working capital requirements. The net cash provided by operating activities for the first quarter of 2011 was $138.2 million compared to $173.3 million for the first quarter of 2010. Cash flow from operating activities

decreased in the first quarter of 2011 compared to the first quarter of 2010 primarily as a result of an increase in cash required to fund changes in trade receivables and accounts payable, partially offset by a decrease in cash used to fund changes in other current assets and other liabilities. In the first quarter of 2011, we paid $15.2 million in connection with the 2010 global settlement with the DOJ regarding our past U.S. sales and marketing practices related to certain therapeutic uses of *Botox*®. In the first quarter of 2011, we paid pension contributions of $3.5 million to our U.S. defined benefit pension plan. We did not make any pension contributions to our U.S. defined benefit pension plan in the first quarter of 2010.

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Net cash provided by investing activities was $479.2 million in the first quarter of 2011 compared to net cash used in investing activities of $80.8 million in the first quarter of 2010. In the first quarter of 2011, we received $649.3 million from the maturities of short-term investments and purchased $149.9 million of short-term investments. Additionally, we invested $17.1 million in new facilities and equipment and $3.3 million in capitalized software. In the first quarter of 2010, we paid $63.7 million, net of cash acquired, for the acquisition of Serica and $1.7 million for a contractual purchase price adjustment related to our 2009 acquisition of Samil Allergan Ophthalmic Joint Venture Company. Additionally, we invested $12.5 million in new facilities and equipment and $2.9 million in capitalized software. We currently expect to invest between $160.0 million and $180.0 million in capital expenditures for manufacturing and administrative facilities, manufacturing equipment and other property, plant and equipment during 2011.

Net cash used in financing activities was $87.9 million in the first quarter of 2011 compared to $42.2 million in the first quarter of 2010. In the first quarter of 2011, we repurchased 2.3 million shares of our common stock for $162.9 million, paid $15.3 million in dividends to stockholders and paid contingent consideration of $2.9 million. This use of cash was partially offset by $8.2 million in net borrowing of notes payable, $80.4 million received from the sale of stock to employees and $4.6 million in excess tax benefits from share-based compensation. In the first quarter of 2010, we repurchased 1.0 million shares of our common stock for $59.6 million, had net repayments of notes payable of $3.5 million and paid $15.1 million in dividends. This use of cash was partially offset by $36.0 million received from the sale of stock to employees.

Effective May 3, 2011, our Board of Directors declared a cash dividend of $0.05 per share, payable June 10, 2011 to stockholders of record on May 20, 2011.

We maintain an evergreen stock repurchase program. Our evergreen stock repurchase program authorizes us to repurchase our common stock for the primary purpose of funding our stock-based benefit plans. Under the stock repurchase program, we may maintain up to 18.4 million repurchased shares in our treasury account at any one time. At March 31, 2011, we held approximately 2.5 million treasury shares under this program. Effective January 1, 2011, our current Rule 10b5-1 plan authorizes our broker to purchase our common stock traded in the open market pursuant to our evergreen stock repurchase program. The terms of the plan set forth a maximum limit of 4.0 million shares to be repurchased, certain quarterly maximum and minimum volume limits, and the plan is cancellable at any time in our sole discretion and in accordance with applicable insider trading laws.

Our 2020 Notes, which were sold at 99.697% of par value with an effective interest rate of 3.41%, are unsecured and pay interest semi-annually on the principal amount of the notes at a rate of 3.375% per annum, and are redeemable at any time at our option, subject to a make-whole provision based on the present value of remaining interest payments at the time of the redemption. The aggregate outstanding principal amount of the 2020 Notes will be due and payable on September 15, 2020, unless earlier redeemed by us.

Our 1.50% Convertible Senior Notes due 2026, or 2026 Convertible Notes, pay interest semi-annually on the principal amount of the notes at a rate of 1.50% per annum and are convertible, at the holder s option, at a conversion rate of 15.7904 shares per $1,000 principal amount of notes if our stock price reaches certain specified thresholds or we call the 2026 Convertible Notes for redemption. We are permitted to redeem the 2026 Convertible Notes at the principal amount plus accrued interest at any time on or after April 5, 2011. Under the terms of the 2026 Convertible Notes, upon conversion we will pay the principal amount in cash and, if the conversion value exceeds the principal amount at the time of conversion, we will also deliver common stock or, at our election, all cash or a combination of cash and common stock for the conversion value in excess of the principal amount.

On March 8, 2011, we announced our intention to redeem the 2026 Convertible Notes at the principal amount plus accrued interest on April 5, 2011. Most note holders have elected to exercise the conversion feature of the 2026 Convertible Notes prior to redemption and we have elected to pay the full conversion value in cash. The conversion value of a note is based on an average of the daily closing price of our stock over an averaging period that commences after we receive a conversion notice from a note holder. We expect to pay approximately $800.0 million in aggregate conversion value in April and May 2011 for the converted notes. We redeemed the notes that were not converted for $8.6 million, representing the aggregate principal amount plus accrued interest, on April 5, 2011.

Our 5.75% Senior Notes due 2016, or 2016 Notes, were sold at 99.717% of par value with an effective interest rate of 5.79%, pay interest semi-annually on the principal amount of the notes at a rate of 5.75% per annum, and are redeemable at any time at our option, subject to a make-whole provision based on the present value of remaining interest payments at the time of the redemption. The aggregate outstanding principal amount of the 2016 Notes is due and payable on April 1, 2016, unless earlier redeemed by us.

At March 31, 2011, we had a committed long-term credit facility, a commercial paper program, a medium-term note program, a shelf registration statement that allows us to issue additional securities, including debt securities, in one or more

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offerings from time to time, a real estate mortgage and various foreign bank facilities. Our committed long-term credit facility expires in May 2012. The termination date can be further extended from time to time upon our request and acceptance by the issuer of the facility for a period of one year from the last scheduled termination date for each request accepted. The committed long-term credit facility allows for borrowings of up to $800.0 million. The commercial paper program also provides for up to $600.0 million in borrowings. Borrowings under the committed long-term credit facility and medium-term note program are subject to certain financial and operating covenants that include, among other provisions, maximum leverage ratios. Certain covenants also limit subsidiary debt. We believe we were in compliance with these covenants at March 31, 2011. At March 31, 2011, we had no borrowings under our committed long-term credit facility, $25.0 million in borrowings outstanding under the medium-term note program, $20.0 million in borrowings outstanding under the real estate mortgage, $36.3 million in borrowings outstanding under various foreign bank facilities and no borrowings under the commercial paper program. Commercial paper, when outstanding, is issued at current short-term interest rates. Additionally, any future borrowings that are outstanding under the long-term credit facility will be subject to a floating interest rate. We may from time to time seek to retire or purchase our outstanding debt.

At December 31, 2010, we had net pension and postretirement benefit obligations totaling $204.7 million. Future funding requirements are subject to change depending on the actual return on net assets in our funded pension plans and changes in actuarial assumptions. In 2011, we expect to pay pension contributions of between $35.0 million and $45.0 million for our U.S. and non-U.S. pension plans and between $1.0 million and $2.0 million for our other postretirement plan.

On January 28, 2011, we entered into a collaboration agreement and a co-promotion agreement with MAP for the exclusive development and commercialization by us and MAP of *Levadex* within the United States to certain headache specialist physicians for the treatment of acute migraine in adults, migraine in adolescents and other indications that may be approved by the parties. Under the terms of the agreements, we made a $60.0 million upfront payment to MAP in February 2011. The terms of the agreements also include up to $97.0 million in additional payments to MAP upon MAP meeting certain development and regulatory milestones.

On May 4, 2011, we announced a license agreement with Molecular Partners AG pursuant to which we obtain exclusive global rights in the field of ophthalmology for MP0112, a Phase II proprietary therapeutic *DARPin*® protein targeting vascular endothelial growth factor receptors under investigation for the treatment of retinal diseases. Under the terms of the agreement, we will make a $45.0 million upfront payment to Molecular Partners AG and potential future milestone and royalty payments.

In May 2011, a generic version of *Elestat*® was launched in the United States and we expect generic versions of *Zymar*® and *Tazorac*® cream to be launched in the United States during 2011. In addition, generic versions of some branded pharmaceutical products sold by our competitors have been launched or are expected to be launched in the United States during 2011. We do not believe that our liquidity will be materially impacted in 2011 by generic competition.

A significant amount of our existing cash and equivalents are held by non-U.S. subsidiaries. We currently plan to use these funds in our operations outside the United States. Withholding and U.S. taxes have not been provided for unremitted earnings of certain non-U.S. subsidiaries because we have reinvested these earnings indefinitely in such operations. At December 31, 2010, we had approximately $2,109.4 million in unremitted earnings outside the United States for which withholding and U.S. taxes were not provided. Tax costs would be incurred if these funds were remitted to the United States.

We believe that the net cash provided by operating activities, supplemented as necessary with borrowings available under our existing credit facilities and existing cash and equivalents and short-term investments, will provide us with sufficient resources to meet our current expected obligations, working capital requirements, debt service, including the conversion of our 2026 Convertible Notes, and other cash needs over the next year.

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ALLERGAN, INC.

Item 3. ***Quantitative and Qualitative Disclosures About Market Risk***

In the normal course of business, our operations are exposed to risks associated with fluctuations in interest rates and foreign currency exchange rates. We address these risks through controlled risk management that includes the use of derivative financial instruments to economically hedge or reduce these exposures. We do not enter into financial instruments for trading or speculative purposes.

To ensure the adequacy and effectiveness of our interest rate and foreign exchange hedge positions, we continually monitor our interest rate swap positions and foreign exchange forward and option positions both on a stand-alone basis and in conjunction with our underlying interest rate and foreign currency exposures, from an accounting and economic perspective.

However, given the inherent limitations of forecasting and the anticipatory nature of the exposures intended to be hedged, we cannot assure you that such programs will offset more than a portion of the adverse financial impact resulting from unfavorable movements in either interest or foreign exchange rates. In addition, the timing of the accounting for recognition of gains and losses related to mark-to-market instruments for any given period may not coincide with the timing of gains and losses related to the underlying economic exposures and, therefore, may adversely affect our consolidated operating results and financial position.

Interest Rate Risk

Our interest income and expense are more sensitive to fluctuations in the general level of U.S. interest rates than to changes in rates in other markets. Changes in U.S. interest rates affect the interest earned on our cash and equivalents and short-term investments and interest expense on our debt, as well as costs associated with foreign currency contracts.

On January 31, 2007, we entered into a nine-year, two-month interest rate swap with a $300.0 million notional amount with semi-annual settlements and quarterly interest rate reset dates. The swap receives interest at a fixed rate of 5.75% and pays interest at a variable interest rate equal to 3-month LIBOR plus 0.368%, and effectively converts $300.0 million of the $800.0 million aggregate principal amount of our 2016 Notes to a variable interest rate. Based on the structure of the hedging relationship, the hedge meets the criteria for using the short-cut method for a fair value hedge. The investment in the derivative and the related long-term debt are recorded at fair value. At March 31, 2011 and December 31, 2010, we recognized in our consolidated balance sheets an asset reported in Investments and other assets and a corresponding increase in Long-term debt associated with the fair value of the derivative of $37.5 million and $42.3 million, respectively. The differential to be paid or received as interest rates change is accrued and recognized as an adjustment of interest expense related to the 2016 Notes. During the first quarter of 2011 and 2010, we recognized $3.8 million, respectively, as a reduction of interest expense due to the differential to be received.

In February 2006, we entered into interest rate swap contracts based on 3-month LIBOR with an aggregate notional amount of $800.0 million, a swap period of 10 years and a starting swap rate of 5.198%. We entered into these swap contracts as a cash flow hedge to effectively fix the future interest rate for our 2016 Notes. In April 2006, we terminated the interest rate swap contracts and received approximately $13.0 million. The total gain is being amortized as a reduction to interest expense over a 10 year period to match the term of the 2016 Notes. As of March 31, 2011, the remaining unrecognized gain, net of tax, of $3.9 million is recorded as a component of accumulated other comprehensive loss.

At March 31, 2011, we had approximately $36.3 million of variable rate debt. If interest rates were to increase or decrease by 1% for the year, annual interest expense, including the effect of the $300.0 million notional amount of the interest rate swap entered into on January 31, 2007, would increase or decrease by approximately $3.4 million. Commercial paper, when outstanding, is issued at current short-term interest rates. Additionally, any future borrowings that are outstanding under the long-term credit facility will be subject to a floating interest rate. Therefore, higher interest costs could occur if interest rates increase in the future.

The following tables present information about certain of our investment portfolio and our debt obligations at March 31, 2011 and December 31, 2010.

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	March 31, 2011 Maturing in							
	2011	2012	2013	2014	2015	Thereafter	Total	Fair Market Value
	(in millions, except interest rates)							
ASSETS								
Cash Equivalents and Short-Term Investments:								
Commercial Paper	$ 1,159.4	$	$	$	$	$	$ 1,159.4	$ 1,159.4
Weighted Average Interest Rate	0.21%						0.21%	
Foreign Time Deposits	227.5						227.5	227.5
Weighted Average Interest Rate	0.47%						0.47%	
Other Cash Equivalents	1,272.1						1,272.1	1,272.1
Weighted Average Interest Rate	0.11%						0.11%	
Total Cash Equivalents and Short-Term Investments	$ 2,659.0	$	$	$	$	$	$ 2,659.0	$ 2,659.0
Weighted Average Interest Rate	0.18%						0.18%	
LIABILITIES								
Debt Obligations:								
Fixed Rate (US$)	$ 648.9	$ 25.0	$	$	$	$ 1,467.0	$ 2,140.9	$ 2,201.7
Weighted Average Interest Rate	5.59%	7.47%				4.74%	5.03%	
Other Variable Rate (non-US$)	36.3						36.3	36.3
Weighted Average Interest Rate	7.19%						7.19%	
Total Debt Obligations (a)	$ 685.2	$ 25.0	$	$	$	$ 1,467.0	$ 2,177.2	$ 2,238.0
Weighted Average Interest Rate	5.67%	7.47%				4.74%	5.06%	
INTEREST RATE DERIVATIVES								
Interest Rate Swaps:								
Fixed to Variable (US$)	$	$	$	$	$	$ 300.0	$ 300.0	$ 37.5
Average Pay Rate						0.67%	0.67%	
Average Receive Rate						5.75%	5.75%	

(a) Total debt obligations in the unaudited condensed consolidated balance sheet at March 31, 2011 include debt obligations of $2,177.2 million and the interest rate swap fair value adjustment of $37.5 million.

	December 31, 2010 Maturing in							
	2011	2012	2013	2014	2015	Thereafter	Total	Fair Market Value
	(in millions, except interest rates)							
ASSETS								
Cash Equivalents and Short-Term Investments:								
Commercial Paper	$ 1,716.0	$	$	$	$	$	$ 1,716.0	$ 1,716.0
Weighted Average Interest Rate	0.25%						0.25%	
Foreign Time Deposits	209.6						209.6	209.6
Weighted Average Interest Rate	0.45%						0.45%	
Other Cash Equivalents	707.0						707.0	707.0
Weighted Average Interest Rate	0.38%						0.38%	
Total Cash Equivalents and Short-Term Investments	$ 2,632.6	$	$	$	$	$	$ 2,632.6	$ 2,632.6
Weighted Average Interest Rate	0.30%						0.30%	
LIABILITIES								
Debt Obligations:								
Fixed Rate (US$)	$ 642.5	$ 25.0	$	$	$	$ 1,466.9	$ 2,134.4	$ 2,221.1
Weighted Average Interest Rate	5.59%	7.47%				4.74%	5.02%	
Other Variable Rate (non-US$)	28.1						28.1	28.1
Weighted Average Interest Rate	6.80%						6.80%	

Total Debt Obligations (a)	$ 670.6	$ 25.0	$	$	$ 1,466.9	$ 2,162.5	$ 2,249.2
Weighted Average Interest Rate	5.64%	7.47%			4.74%	5.05%	
INTEREST RATE DERIVATIVES							
Interest Rate Swaps:							
Fixed to Variable (US$)	$	$	$	$	$ 300.0	$ 300.0	$ 42.3
Average Pay Rate					0.67%	0.67%	
Average Receive Rate					5.75%	5.75%	

(a) Total debt obligations in the unaudited condensed consolidated balance sheet at December 31, 2010 include debt obligations of $2,162.5 million and the interest rate swap fair value adjustment of $42.3 million.

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Foreign Currency Risk

Overall, we are a net recipient of currencies other than the U.S. dollar and, as such, benefit from a weaker dollar and are adversely affected by a stronger dollar relative to major currencies worldwide. Accordingly, changes in exchange rates, and in particular a strengthening of the U.S. dollar, may negatively affect our consolidated revenues or operating costs and expenses as expressed in U.S. dollars.

From time to time, we enter into foreign currency option and forward contracts to reduce earnings and cash flow volatility associated with foreign exchange rate changes to allow our management to focus its attention on our core business issues. Accordingly, we enter into various contracts which change in value as foreign exchange rates change to economically offset the effect of changes in the value of foreign currency assets and liabilities, commitments and anticipated foreign currency denominated sales and operating expenses. We enter into foreign currency option and forward contracts in amounts between minimum and maximum anticipated foreign exchange exposures, generally for periods not to exceed 18 months.

We use foreign currency option contracts, which provide for the sale or purchase of foreign currencies to offset foreign currency exposures expected to arise in the normal course of our business. While these instruments are subject to fluctuations in value, such fluctuations are anticipated to offset changes in the value of the underlying exposures.

All of our outstanding foreign currency option contracts are entered into to reduce the volatility of earnings generated in currencies other than the U.S. dollar, primarily earnings denominated in the Canadian dollar, Mexican peso, Australian dollar, Brazilian real, euro, Korean won and Turkish lira. Current changes in the fair value of open foreign currency option contracts and any realized gains (losses) on settled contracts are recorded through earnings as Other, net in the accompanying unaudited condensed consolidated statements of earnings. The premium costs of purchased foreign exchange option contracts are recorded in Other current assets and amortized to Other, net over the life of the options.

All of our outstanding foreign exchange forward contracts are entered into to offset the change in value of certain intercompany receivables or payables that are subject to fluctuations in foreign currency exchange rates. The realized and unrealized gains and losses from foreign currency forward contracts and the revaluation of the foreign denominated intercompany receivables or payables are recorded through Other, net in the accompanying unaudited condensed consolidated statements of earnings.

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The following table provides information about our foreign currency derivative financial instruments outstanding as of March 31, 2011 and December 31, 2010. The information is provided in U.S. dollars, as presented in our unaudited condensed consolidated financial statements:

	March 31, 2011		December 31, 2010	
	Notional Amount (in millions)	Average Contract Rate or Strike Amount	Notional Amount (in millions)	Average Contract Rate or Strike Amount
Foreign currency forward contracts:				
(Receive U.S. dollar/pay foreign currency)				
Euro	$ 1.4	1.39	$	
Japanese yen	7.4	80.84	6.0	84.09
Australian dollar	20.5	0.98	15.7	0.98
New Zealand dollar	1.5	0.73	1.1	0.74
Poland zloty	1.6	2.93	2.8	3.03
Singapore dollar	3.9	1.28		
	$ 36.3		$ 25.6	
Estimated fair value	$ (0.9)		$ (0.9)	
Foreign currency forward contracts:				
(Pay U.S. dollar/receive foreign currency)				
Euro	$ 41.7	1.39	$ 39.9	1.33
Estimated fair value	$ 0.7		$ 0.2	
Foreign currency sold put options:				
Canadian dollar	$ 71.6	1.03	$ 68.1	1.04
Mexican peso	15.3	12.79	20.0	12.73
Australian dollar	45.2	0.88	44.2	0.87
Brazilian real	38.3	1.92	36.9	1.92
Euro	144.3	1.34	139.4	1.34
Korean won	13.2	1153.66	17.3	1153.22
Turkish lira	16.5	1.56	20.5	1.55
	$ 344.4		$ 346.4	
Estimated fair value	$ 4.6		$ 10.4	

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ALLERGAN, INC.

Item 4. ***Controls and Procedures***

Evaluation of Disclosure Controls and Procedures

We maintain disclosure controls and procedures that are designed to ensure that information required to be disclosed in our Exchange Act reports is recorded, processed, summarized and reported within the time periods specified in the Securities and Exchange Commission s rules and forms, and that such information is accumulated and communicated to our management, including our Principal Executive Officer and our Principal Financial Officer, as appropriate, to allow timely decisions regarding required disclosures. Our management, including our Principal Executive Officer and our Principal Financial Officer, does not expect that our disclosure controls or procedures will prevent all error and all fraud. A control system, no matter how well conceived and operated, can provide only reasonable, not absolute, assurance that the objectives of the control system are met. Further, the benefits of controls must be considered relative to their costs. Because of the inherent limitations in all control systems, no evaluation of controls can provide absolute assurance that all control issues and instances of fraud, if any, have been detected. These inherent limitations include the realities that judgments in decision-making can be faulty, and that breakdowns can occur because of simple error or mistake. Additionally, controls can be circumvented by the individual acts of some persons, by collusion of two or more people, or by management override of the control. The design of any system of controls is also based in part upon certain assumptions about the likelihood of future events, and there can be no assurance that any design will succeed in achieving its stated goals under all potential future conditions. Because of the inherent limitations in a cost-effective control system, misstatements due to error or fraud may occur and not be detected. Also, we have investments in certain unconsolidated entities. As we do not control or manage these entities, our disclosure controls and procedures with respect to such entities are necessarily substantially more limited than those we maintain with respect to our consolidated subsidiaries.

We carried out an evaluation, under the supervision and with the participation of our management, including our Principal Executive Officer and our Principal Financial Officer, of the effectiveness of the design and operation of our disclosure controls and procedures as of March 31, 2011, the end of the quarterly period covered by this report. Based on the foregoing, our Principal Executive Officer and our Principal Financial Officer concluded that, as of the end of the period covered by this report, our disclosure controls and procedures were effective and were operating at the reasonable assurance level.

Further, management determined that, as of March 31, 2011, there were no changes in our internal control over financial reporting that occurred during the quarterly period covered by this report that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

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ALLERGAN, INC.

PART II OTHER INFORMATION

Item 1. ***Legal Proceedings***

The following supplements and amends the discussion set forth under Part I, Item 3 Legal Proceedings of our Annual Report on Form 10-K for the year ended December 31, 2010.

Clayworth v. Allergan, et al.

In August 2004, James Clayworth, R.Ph., doing business as Clayworth Pharmacy, filed a complaint entitled Clayworth v. Allergan, *et al.* in the Superior Court of the State of California for the County of Alameda. The complaint, as amended, named us and 12 other defendants and alleged unfair business practices, including a price fixing conspiracy relating to the reimportation of pharmaceuticals from Canada. The complaint sought damages, equitable relief, attorneys fees and costs. In January 2007, the court entered a notice of entry of judgment of dismissal against the plaintiffs, dismissing the plaintiffs complaint. On the same date, the plaintiffs filed a notice of appeal with the Court of Appeal of the State of California. In July 2008, the court of appeal affirmed the superior court s ruling, granting our motion for summary judgment. In August 2008, the plaintiffs filed a petition for rehearing with the court of appeal, which the court denied. In September 2008, the plaintiffs filed a petition for review with the Supreme Court of the State of California, which the supreme court granted in November 2008. In July 2010, the supreme court reversed the court of appeal s judgment and remanded the case to the superior court for further proceedings. In October 2010, plaintiffs filed a challenge to the assignment of this matter to the presiding judge alleging a conflict of interest. In November 2010, plaintiffs challenge was denied. In December 2010, plaintiffs filed a petition for writ of mandate in the Court of Appeal of the State of California seeking to overturn the order denying their challenge. In December 2010, the court of appeal denied the petition. In December 2010, plaintiffs filed a petition for review with the Supreme Court of the State of California. In January 2011, the court set trial for August 1, 2011. In February 2011, the supreme court denied plaintiffs petition for review. In March 2011, the court entered judgment in favor of defendants pursuant to orders granting motions for summary judgment. In April 2011, plaintiffs filed a notice of appeal to the Court of Appeal of the State of California.

Allergan, Inc. v. Cayman Chemical Company, et al.

In November 2007, we filed a complaint captioned Allergan, Inc. v. Cayman Chemical Company, Jan Marini Skin Research, Inc., Athena Cosmetics, Inc., Dermaquest, Inc., Intuit Beauty, Inc., Civic Center Pharmacy and Photomedex, Inc. in the U.S. District Court for the Central District of California. In the complaint, we allege that the defendants are infringing U.S. Patent No. 6,262,105 licensed to us by Murray A. Johnstone, M.D. In March 2008, we filed a second amended complaint to add Dr. Johnstone, the holder of U.S. Patent No. 6,262,105, as a plaintiff and to add Global MDRx and ProCyte Corporation, or ProCyte, as defendants. In April 2008, we filed a motion for leave to file a third amended complaint to add patent infringement claims relating to U.S. Patent No. 7,351,404 against the defendants, and to add Athena Bioscience, LLC and Cosmetic Alchemy, LLC as additional defendants.

In 2008, we entered into settlement agreements with Jan Marini Skin Research, Inc., Intuit Beauty, Inc., Photomedex, Inc. and ProCyte pursuant to which each party agreed to acknowledge the validity of the patents in exchange for dismissing all claims against such defendant. In July 2008, the clerk of the court entered a default judgment against Global MDRx for failure to defend against the summons. In August 2008, the court dismissed Intuit Beauty, Inc. and Jan Marini Skin Research, Inc. with prejudice. In September 2008, we and Cayman Chemical Company entered into a settlement agreement under which Cayman Chemical Company agreed to cease selling certain compounds to be used in particular types of products in exchange for dismissing all claims against them. In December 2008, we entered into a settlement agreement with Athena Bioscience, LLC under which they agreed to cease selling certain products and acknowledged the validity of our patents in exchange for our dismissing all claims against them.

In January 2009, we, along with Dr. Johnstone, filed a motion for leave to file a fourth amended complaint adding Pharma Tech, Inc., Dimensional Merchandising, Inc. and Cosmetic Technologies, Inc. as new defendants. In February 2009, we, along with Dr. Johnstone, filed a motion for default judgment and injunction against Global MDRx and the court granted our motion. In April 2009, we and Cosmetic Technologies, Inc. entered into a settlement agreement under which Cosmetic Technologies, Inc. agreed to cease manufacturing and selling certain products and acknowledge the validity of our patents in exchange for our dismissing all claims against them.

In March 2009, we filed a complaint captioned Allergan, Inc.; Murray A Johnstone, M.D.; and Duke University v. Athena Cosmetics, Inc.; Cosmetic Alchemy, LLC; Northwest Cosmetic Laboratories, LLC; Pharma Tech International, Inc.; Dimensional Merchandising, Inc.; Stella

International, LLC; Product Innovations, LLC; Metrics, LLC; Nutra-Luxe M.D.,

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LLC; Skin Research Laboratories, Inc.; Lifetech Resources LLC; Rocasuba, Inc.; Peter Thomas Roth Labs LLC; and Peter Thomas Roth, Inc. in the U.S. District Court for the Central District of California alleging infringement of U.S. Patent Nos. 6,262,105, 7,351,404, and 7,388,029. In June 2009, we and defendants La Canada Ventures, Inc. and Susan Lin, M.D. entered into a settlement agreement under which La Canada Ventures, Inc. and Susan Lin, M.D. agreed to cease manufacturing and selling certain products and acknowledge the validity of our patents in exchange for our dismissing all claims against La Canada Ventures, Inc. and Susan Lin, M.D.

In June 2009, the court consolidated Allergan, Inc.; Murray A Johnstone, M.D.; and Duke University v. Athena Cosmetics, Inc., *et al.* with Allergan, Inc. v. Cayman Chemical Company, *et al.* and set an October 2010 trial date for both cases. In October 2009, the defendants filed answers, amended answers and/or counterclaims to our first amended complaint. In February 2010, we and Athena Cosmetic, Inc. filed a stipulation with the court to bifurcate Athena Cosmetic, Inc. s antitrust and Lanham Act counterclaims into separate trials. In February 2010, Athena Cosmetic, Inc., Pharma Tech and Northwest Cosmetic filed a motion for judgment on the pleadings regarding our claim for violation of the California unfair competition statute. In March 2010, the court granted Athena Cosmetic, Inc., Pharma Tech and Northwest Cosmetic s motion for judgment on the pleadings. In May 2010, we entered into a settlement agreement with Nutra-Luxe M.D., LLC, under which Nutra-Luxe M.D., LLC agreed to cease manufacturing and selling certain products and acknowledge the validity of our patents in exchange for our dismissing all claims against them. In May 2010, pursuant to a stipulation filed by the plaintiffs and all defendants against whom there are currently claims pending in the two consolidated actions, the court entered an order stating that a final judgment will be entered on the dismissal of our unfair competition claim against the defendants, permitting us to appeal the dismissal without further delay to the U.S. Court of Appeals for the Federal Circuit, and further stating that all U.S. District Court proceedings in both consolidated actions will be stayed pending completion of our appeal of the dismissal of our unfair competition claim. In May 2010, we filed a notice of appeal with the court of appeals. On March 9, 2011, the court of appeals heard oral argument on the appeal and took the matter under submission.

Kramer et al. v. Allergan, Inc.

In July 2008, a complaint entitled Kramer, Bryant, Spears, Doolittle, Clark, Whidden, Powell, Moore, Hennessey, Sody, Breeding, Downey, Underwood-Boswell, Reed-Momot, Purdon & Hahn v. Allergan, Inc. was filed in the Superior Court for the State of California for the County of Orange. The complaint makes allegations against us relating to *Botox*® and *Botox*® Cosmetic including failure to warn, manufacturing defects, negligence, breach of implied and express warranties, deceit by concealment and negligent misrepresentation and seeks damages, attorneys fees and costs. In 2009, the plaintiffs Hennessey, Hahn, Underwood-Boswell, Purdon, Moore, Clark, Reed-Momot and Whidden were dismissed without prejudice. The trial related to plaintiff Spears began in January 2010. In March 2010, the jury returned a verdict in our favor and the court entered a judgment on the special verdict. In April 2010, plaintiff Spears filed a motion for a new trial, which the court denied in May 2010. In June 2010, we and plaintiff Spears entered into a settlement agreement under which we agreed to waive costs in exchange for plaintiff Spears agreeing not to appeal the judgment. In September 2010, the trial related to plaintiff Bryant began and we subsequently entered into a settlement agreement with plaintiff Bryant. In April 2011, the court set the trial related to plaintiff Doolittle for November 7, 2011 or, in the alternative, for January 17, 2012.

Zymar® Patent Litigation

In October 2007, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Apotex, indicating that Apotex had filed an ANDA with the FDA for a generic version of *Zymar*®. In the certification, Apotex contends that U.S. Patent Nos. 5,880,283 and 6,333,045, or the 045 patent, both of which are licensed to us and are listed in the Orange Book under *Zymar*®, are invalid and/or not infringed by the proposed Apotex product. In November 2007, we, Senju Pharmaceutical Co., Ltd., or Senju, and Kyorin Pharmaceutical Co., Ltd., or Kyorin, filed a complaint captioned Allergan, Inc., Senju Pharmaceutical Co., Ltd. and Kyorin Pharmaceutical Co., Ltd. v. Apotex Inc., *et al.* in the U.S. District Court for the District of Delaware. The complaint alleges infringement of the 045 patent. In January 2008, Apotex filed an answer and a counterclaim, as well as a motion to partially dismiss the plaintiffs complaint. In February 2008, we, Senju and Kyorin filed a response of non-opposition to Apotex s motion to partially dismiss the complaint. A three-day bench trial was conducted in January 2010. In June 2010, the court ruled that Apotex s proposed generic version of *Zymar*® infringes claims 1-3, 6, 7 and 9 of the 045 patent and that claims 1-3 and 6-9 are invalid as obvious. The court further ruled that Apotex failed to prove that claims 6 and 7 are invalid for lack of enablement and that Apotex failed to prove that the 045 patent is unenforceable for inequitable conduct. In June 2010, we, Senju and Kyorin filed a motion for a new trial or, alternatively, to amend judgment and findings regarding claim 7. In July 2010, Apotex filed an answer to our motion and we filed a reply to Apotex s answer to our motion. In November 2010, the court dismissed our motion for a new trial without prejudice to renew and opened the record of the litigation so that additional evidence may be submitted. In April 2011, the court held an evidentiary hearing and another evidentiary hearing is scheduled for May 2011.

In August 2010, we filed a statement of claim entitled Allergan, *et al.* & Kyorin Pharmaceutical Co., LTD v. Apotex

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Inc., *et al.* in the Federal Court of Canada at Ottawa, Ontario, Canada. The statement of claim alleges that Apotex s product infringes Canadian Patent No. 1,340,316 covering *Zymar®*. In September 2010, Apotex filed a motion to strike the statement of claim. In November 2010, the court dismissed the motion to strike. In November 2010, Apotex filed a notice of appeal regarding the dismissed motion to strike. In April 2011, the court of appeal heard oral argument on the appeal and dismissed the appeal.

In April 2011, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Lupin Limited, or Lupin, indicating that Lupin had filed an ANDA with the FDA seeking approval of a generic form of *Zymar®* gatifloxacin 0.3% ophthalmic solution. In the certification, Lupin contends that U.S. Patent Nos. 5,880,283 and the 405 patent, listed in the Orange Book under *Zymar®*, are invalid and/or not infringed by the proposed Lupin product.

Combigan® Patent Litigation

In February 2009 and April 2009, we received paragraph 4 invalidity and noninfringement Hatch-Waxman Act certifications from Sandoz, Inc., or Sandoz, and Hi-Tech Pharmacal Co. Inc., or Hi-Tech, respectively, indicating that Sandoz and Hi-Tech had filed ANDAs seeking approval of generic forms of *Combigan®*, a brimonidine tartrate 0.2%, timolol 0.5% ophthalmic solution. In their separate certifications, Sandoz and Hi-Tech each contend that U.S. Patent Nos. 7,030,149 and 7,320,976, listed in the Orange Book under *Combigan®*, are invalid and/or not infringed by the proposed Sandoz product and by the proposed Hi-Tech product. We filed complaints against Sandoz and Hi-Tech in the U.S. District Court for the Eastern District of Texas in April 2009 and June 2009, respectively, alleging, in each case, that the defendant s proposed product infringes U.S. Patent Nos. 7,030,149 and 7,320,976. In October 2009, we filed a motion to consolidate the Hi-Tech action and the Sandoz action and the court granted our motion to consolidate the two actions.

In September 2009, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Alcon Research, Ltd., or Alcon, indicating that Alcon had filed an ANDA seeking approval of a generic version of *Combigan®*. In the certification, Alcon contends that U.S. Patent Nos. 7,030,149, 7,320,976 and 7,323,463, listed in the Orange Book under *Combigan®*, are invalid and/or not infringed by the proposed Alcon product. In November 2009, we filed a complaint against Alcon in the U.S. District Court for the Eastern District of Texas, Marshall Division. The complaint alleges that Alcon s proposed product infringes U.S. Patent Nos. 7,030,149, 7,320,976 and 7,323,463.

In October 2009 and November 2009, we received amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certifications from Sandoz and Hi-Tech, respectively, indicating that Sandoz and Hi-Tech had filed ANDAs seeking approval of generic forms of *Combigan®*. In their separate certifications, Sandoz and Hi-Tech each contend that U.S. Patent No. 7,323,463, listed in the Orange Book under *Combigan®*, is invalid and/or not infringed by the proposed Sandoz and Hi-Tech products. In November 2009, we filed an amended complaint against Sandoz and Hi-Tech for patent infringement to assert U.S. Patent No. 7,323,463. Sandoz filed an answer and counterclaims to our amended complaint in November 2009 and Hi-Tech filed an answer and counterclaims in December 2009. We filed an answer to Sandoz s counterclaims in December 2009 and an answer to Hi-Tech s counterclaims in January 2010. In January 2010, the Hi-Tech action and the Sandoz action were consolidated with the Alcon action.

In February 2010, we received amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certifications from Sandoz and Hi-Tech indicating that Sandoz and Hi-Tech had filed ANDAs seeking approval of generic forms of *Combigan®*. In their separate certifications, Sandoz and Hi-Tech contend that U.S. Patent No. 7,642,258, listed in the Orange Book under *Combigan®*, is invalid and/or not infringed by the proposed Sandoz and Hi-Tech products. In March 2010, we filed a second amended complaint against Sandoz and Hi-Tech for patent infringement to assert U.S. Patent No. 7,642,258. Hi-Tech and Sandoz filed an answer and counterclaims to our second amended complaint in March 2010 and April 2010, respectively. In April 2010, we filed answers to Hi-Tech and Sandoz s counterclaims. In April 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Alcon indicating that Alcon had filed an ANDA seeking approval of a generic form of *Combigan®*. In their certification, Alcon contends that U.S. Patent No. 7,642,258, listed in the Orange Book under *Combigan®*, is invalid and/or not infringed by the proposed Alcon product. In April 2010, we filed a first amended complaint against Alcon for patent infringement to assert U.S. Patent No. 7,642,258. In May 2010, Alcon filed an answer and counterclaims to our first amended complaint. In June 2010, we filed an answer to Alcon s counterclaims. The court has scheduled an August 1, 2011 trial date for the consolidated Hi-Tech, Sandoz and Alcon actions.

In May 2010, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Apotex Corp. and Apotex indicating that Apotex had filed an ANDA seeking approval of a generic version of *Combigan®*. In the certification, Apotex contends that U.S. Patent Nos. 7,030,149, 7,320,976, 7,323,463 and 7,642,258 listed in the Orange Book under *Combigan®*, are invalid and/or not infringed by the proposed Apotex product. In June 2010, we filed a complaint against Apotex in the U.S. District Court for the Eastern District of Texas, Marshall Division. The complaint alleges that Apotex s proposed product infringes U.S. Patent Nos. 7,030,149, 7,320,976, 7,323,463 and 7,642,258. In June 2010, we filed

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an amended complaint. In July 2010, Apotex filed an answer and counterclaims to our first amended complaint. In August 2010, we filed an answer to Apotex s counterclaims. In September 2010, the Hi-Tech action, the Sandoz action, and the Alcon action were consolidated with the Apotex action.

In July 2010, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Watson Laboratories, Inc., Watson Pharma, Inc. and Watson Pharmaceuticals, Inc., or Watson, indicating that Watson had filed an ANDA seeking approval of a generic version of *Combigan®*. In the certification, Watson contends that U.S. Patent Nos. 7,030,149, 7,320,976, 7,323,463 and 7,642,258, listed in the Orange Book under *Combigan®*, are invalid and/or not infringed by the proposed Watson product. In September 2010, we filed a complaint against Watson in the U.S. District Court for the Eastern District of Texas, Marshall Division. The complaint alleges that Watson s proposed product infringes U.S. Patent Nos. 7,030,149, 7,320,976, 7,323,463 and 7,642,258. In October 2010, Watson filed an unopposed motion to dismiss without prejudice Watson Pharmaceuticals, Inc. and Watson Pharma, Inc., which the court granted. In October 2010, Watson filed an answer to the complaint and counterclaims. In November 2010, we filed an answer to Watson s counterclaims. In March 2011, the Hi-Tech action, the Sandoz action, the Alcon action, and the Apotex action were consolidated with the Watson action. In April 2011, the court issued its Markman ruling.

In December 2009, we received a Notice of Allegation letter from Sandoz Canada Inc., or Sandoz Canada, indicating that Sandoz Canada had filed an Abbreviated New Drug Submission, or ANDS, under paragraphs 5(1)(b)(iii), 5(1)(b)(iv) and 5(3) of the Patented Medicines (Notice of Compliance) Regulations for approval of a generic version of *Combigan®* (DIN 02248347). In the letter, Sandoz Canada contends that Canadian Patent Nos. 2,173,974, 2,225,626 and 2,440,764 are invalid and/or not infringed by the proposed Sandoz Canada product. In February 2010, we filed a notice of application in the Canadian Federal Court. The application alleges that Sandoz Canada s proposed product infringes Canadian Patent Nos. 2,225,626 and 2,440,764. In February 2010, we received a Notice of Allegation letter from Sandoz Canada indicating that Sandoz Canada had filed an ANDS under paragraphs 5(1)(b)(iii), 5(1)(b)(iv) and 5(3) of the Patented Medicines (Notice of Compliance) Regulations for approval of a generic version of *Combigan®*. In the letter, Sandoz Canada contends that Canadian Patent No. 2,357,014 is invalid and/or not infringed by the proposed Sandoz Canada product. In March 2010, we filed a notice of application in the Canadian Federal Court. The application alleges that Sandoz Canada s proposed product infringes Canadian Patent No. 2,357,014. In May 2010, Sandoz Canada filed two motions to strike the application regarding Canadian Patent No. 2,225,626. In June 2010, the court denied Sandoz Canada s first motion to strike. In August 2010, we entered into an agreement to discontinue our notice of application relating to Canadian Patent No. 2,357,014 in exchange for Sandoz Canada s withdrawing its pending motion to strike the application regarding Canadian Patent No. 2,225,626. In April 2011, the court set the trial in this case for October 17, 2011.

In August 2010, we received a Notice of Allegation letter from Apotex Canada Inc., or Apotex Canada, indicating that Apotex Canada had filed an ANDS under paragraphs 5(1)(b)(iii), 5(1)(b)(iv) and 5(3) of the Patented Medicines (Notice of Compliance) Regulations for approval of a generic version of *Combigan®* (DIN 02248347). In the letter, Apotex Canada contends that Canadian Patent Nos. 2,173,974, 2,225,626, 2,357,014 and 2,440,764 are invalid and/or not infringed by the proposed Apotex Canada product. In September 2010, we filed a notice of application in the Canadian Federal Court. The application alleges that Apotex Canada s proposed product infringes Canadian Patent Nos. 2,225,626, 2,357,014 and 2,440,764. In December 2010, the court set the trial in this case for November 28, 2011.

Sanctura XR® Patent Litigation

In June 2009, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Watson, through its subsidiary Watson Laboratories, Inc. Florida, indicating that Watson had filed an ANDA seeking approval of a generic form of *Sanctura XR®*, trospium 60 mg. chloride extended release capsules. In the certification, Watson contends that U.S. Patent No. 7,410,978, listed in the Orange Book under *Sanctura XR®*, is invalid and/or not infringed by the proposed Watson product.

In July 2009, we, Endo Pharmaceuticals Solutions, Inc., or Endo, and Supernus Pharmaceuticals, Inc., or Supernus, filed a complaint against Watson, Watson Laboratories, Inc. Florida, and Watson Pharma, Inc. in the U.S. District Court for the District of Delaware. The complaint alleges that Watson s proposed product infringes U.S. Patent No. 7,410,978. In August 2009, Watson filed an answer and counterclaims to our complaint. In September 2009, we filed an answer to Watson s counterclaims. In July 2010, Watson filed an amended and supplemental answer and counterclaims to our complaint. In August 2010, we filed an answer to Watson s counterclaims.

In November 2009, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Sandoz indicating that Sandoz had filed an ANDA seeking approval of a generic form of *Sanctura XR®*, trospium 60 mg. chloride extended release capsules. In the certification, Sandoz contends that U.S. Patent No. 7,410,978, listed in the Orange Book under *Sanctura XR®*, is invalid and/or not infringed by the proposed Sandoz product. In November 2009, we, Endo and Supernus filed a complaint against Sandoz in the U.S. District Court for the District of Delaware. The complaint alleges that

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Sandoz s proposed product infringes U.S. Patent No. 7,410,978. In January 2010, Sandoz filed an answer and counterclaims to our complaint. In February 2010, we filed an answer to Sandoz s counterclaims. In March 2010, the court consolidated the Watson and Sandoz actions and scheduled a trial date for May 2, 2011.

In April 2010, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Paddock Laboratories, Inc., or Paddock, indicating that Paddock had filed an ANDA seeking approval of a generic form of *Sanctura XR®*, trospium 60 mg. chloride extended release capsules. In the certification, Paddock contends that U.S. Patent No. 7,410,978, listed in the Orange Book under *Sanctura XR®*, is invalid and/or not infringed by the proposed Paddock product. In June 2010, we, Endo and Supernus filed a complaint against Paddock in the U.S. District Court for the District of Delaware. The complaint alleges that Paddock s proposed product infringes U.S. Patent No. 7,410,978. In July 2010, Paddock filed an answer and counterclaims to our complaint. In August 2010, we filed an answer to Paddock s counterclaims.

In August 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Paddock indicating that Paddock had filed an ANDA seeking approval of a generic form of *Sanctura XR®*. In their certification, Paddock contends that U.S. Patent Nos. 7,759,359 and 7,763,635, listed in the Orange Book under *Sanctura XR®*, are invalid and/or not infringed by the proposed Paddock product. In August 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Watson indicating that Watson had filed an ANDA seeking approval of a generic form of *Sanctura XR®*. In their certification, Watson contends that U.S. Patent Nos. 7,759,359 and 7,763,635, listed in the Orange Book under *Sanctura XR®*, are invalid and/or not infringed by the proposed Watson product.

In September 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Watson indicating that Watson had filed an ANDA seeking approval of a generic form of *Sanctura XR®*. In their certification, Watson contends that U.S. Patent Nos. 7,781,448 and 7,781,449, listed in the Orange Book under *Sanctura XR®*, are invalid and/or not infringed by the proposed Watson product. In September 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Paddock indicating that Paddock had filed an ANDA seeking approval of a generic form of *Sanctura XR®*. In their certification, Paddock contends that U.S. Patent Nos. 7,781,448 and 7,781,449 listed in the Orange Book under *Sanctura XR®*, are invalid and/or not infringed by the proposed Paddock product. In September 2010, the court consolidated the Watson and Sandoz action with the Paddock action.

In October 2010, we, Endo and Supernus filed a complaint against Watson in the U.S. District Court for the District of Delaware. The complaint alleges that Watson s proposed product infringes U.S. Patent Nos. 7,781,448 and 7,781,449. In October 2010, Watson filed an answer and counterclaims in response to the complaint. In October 2010, we, Endo and Supernus filed a complaint against Paddock in the U.S. District Court for the District of Delaware. The complaint alleges that Paddock s proposed product infringes U.S. Patent Nos. 7,781,448 and 7,781,449. In October 2010 and November 2010, Paddock filed answers and counterclaims in response to the complaints. In October 2010, we, Endo and Supernus filed a complaint against Watson and an amended complaint against Paddock and another defendant in the United States District Court for the District of Delaware. The complaint and amended complaint allege that the defendants proposed products infringe U.S. Patent Nos. 7,781,448 and 7,781,449. In October 2010, Paddock filed an answer to the first amended complaint and counterclaims regarding U.S. Patent No. 7,410,978.

In November 2010, Paddock filed an answer to the amended complaint and counterclaims regarding U.S. Patent Nos. 7,781,448 and 7,781,449. In November 2010, we received an amended paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Sandoz indicating that Sandoz had filed an ANDA seeking approval of a generic form of *Sanctura XR®*, trospium 60 mg. chloride extended release capsules. In their certification, Sandoz contends that U.S. Patent Nos. 7,759,359, 7,763,635, 7,781,448 and 7,781,449, listed in the Orange Book under *Sanctura XR®*, are invalid and/or not infringed by the proposed Sandoz product.

In December 2010, we, Endo, and Supernus filed an answer to Paddock s counterclaims with respect to U.S. Patent Nos. 7,410,978, 7,781,448, and 7,781,449. In December 2010, we, Endo, and Supernus filed an answer to Watson s counterclaims with respect to U.S. Patent Nos. 7,781,448 and 7,781,449. In December 2010, we, Endo, and Supernus filed an amended answer to Paddock s counterclaims with respect to U.S. Patent Nos. 7,410,978, 7,781,448, and 7,781,449, and brought an infringement claim regarding U.S. Patent No. 7,759,359. In December 2010, we, Endo, and Supernus filed an amended answer to Watson s counterclaims with respect to U.S. Patent Nos. 7,410,978, 7,781,448 and 7,781,449, and brought an infringement claim regarding U.S. Patent No. 7,759,359.

In January 2011, we, Endo, and Supernus filed a complaint against Sandoz in the United States District Court for the District of Delaware. The complaint alleges that Sandoz s proposed product infringes U.S. Patent Nos. 7,759,359, 7,763,635,

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7,781,448 and 7,781,449. In February 2011, Sandoz filed an answer to our complaint and counterclaims. In February 2011, the court consolidated this action with the Watson, Sandoz, and Paddock actions.

In March 2011, Watson filed an answer to our complaint and counterclaims regarding U.S. Patent Nos. 7,781,448 and 7,781,449. In April 2011, we, Endo, and Supernus filed an amended answer to Watson s counterclaims with respect to U.S. Patent Nos. 7,781,448 and 7,781,449.

Latisse® Patent Litigation

In July 2010, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Apotex indicating that Apotex had filed an ANDA seeking approval of a generic form of *Latisse®*, a bimatoprost 0.3% ophthalmic solution. In the certification, Apotex contends that U.S. Patent Nos. 7,351,404 and 7,388,029, listed in the Orange Book under *Latisse®*, are invalid and/or not infringed by the proposed Apotex product. In September 2010, we and Duke University filed a complaint against Apotex in the U.S. District Court for the Middle District of North Carolina. The complaint alleges that Apotex s proposed product infringes U.S. Patent Nos. 7,351,404, 7,388,029 and 6,403,649. In November 2010, Apotex filed an answer to the complaint and counterclaims. In January 2011, we filed an answer to Apotex s counterclaims.

In March 2011, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Sandoz indicating that Sandoz had filed an ANDA seeking approval of a generic form of *Latisse®*, a bimatoprost 0.3% ophthalmic solution. In the certification, Sandoz contends that U.S. Patent Nos. 7,351,404 and 7,388,029, listed in the Orange Book under *Latisse®*, are invalid and/or not infringed by the proposed Sandoz product. In April 2011, we and Duke University filed a complaint against Sandoz in the U.S. District Court for the Middle District of North Carolina. The complaint alleges that Sandoz s proposed product infringes U.S. Patent Nos. 7,351,404, 7,388,029 and 6,403,649.

Zymaxid® Patent Litigation

In February 2011, we received a paragraph 4 invalidity and noninfringement Hatch-Waxman Act certification from Lupin, indicating that Lupin had filed an ANDA with the FDA seeking approval of a generic form of *Zymaxid®* gatifloxacin 0.05% ophthalmic solution. In the certification, Lupin contends that U.S. Patent Nos. 5,880,283 and 6,333,045, listed in the Orange Book under *Zymaxid®*, are invalid and/or not infringed by the proposed Lupin product. In March 2011, we, Senju and Kyorin filed a complaint captioned Senju Pharmaceutical Co., Ltd., Kyorin Pharmaceutical Co., Ltd., and Allergan, Inc. v. Lupin Limited and Lupin Pharmaceuticals, Inc. in the U.S. District Court for the District of Delaware. The complaint alleges that Lupin s proposed product infringes U.S. Patent Nos. 5,880,283 and 6,333,045.

Government Investigations

In June 2010, we received service of process of a Subpoena from the U.S. Securities and Exchange Commission, or SEC. The subpoena requests the production of documents relating to our affiliation with Acadia Pharmaceuticals, Inc., or Acadia, and our sale of Acadia securities. In September 2010, we produced documents responsive to the Subpoena. In January and March 2011, the SEC issued additional Subpoenas seeking further information, which has been provided.

Stockholder Derivative Litigation

In November 2010, we received a demand for inspection of books and records from U.F.C.W. Local 1776 & Participating Employers Pension Fund, or U.F.C.W. In November 2010, U.F.C.W. filed a motion to intervene in the Louisiana Municipal Police Employees Retirement System action, which was denied by the court in January 2011. In February 2011, U.F.C.W. filed a complaint to compel inspection of books and records in the Court of Chancery of the State of Delaware. In March 2011, we filed an answer to the complaint and the court scheduled the final hearing for April 27, 2011. In April 2011, the court ordered that we produce a limited number of documents to the court for in camera inspection.

In September 2010, Pompano Beach Police & Firefighters Retirement System and Western Washington Laborers-Employers Pension Trust filed a stockholder derivative complaint against our then-current Board of Directors and Allergan, Inc. in the U.S. District Court for the Central District of California. The complaint alleges violations of federal securities laws, breaches of fiduciary duties, abuse of control, gross mismanagement, and corporate waste and seeks, among other things, damages, corporate governance reforms, attorneys fees, and costs. In September 2010, plaintiffs filed a motion for consolidation with the Himmel and Rosenbloom actions, which the court granted in October 2010. In November 2010, the plaintiffs filed their consolidated complaint. In December 2010, we filed a motion to stay the consolidated action in favor of the Louisiana Municipal Police Employees Retirement System action. In December 2010, we and the individual defendants filed motions to dismiss the consolidated complaint. In March 2011, the court denied the motion to stay the consolidated action and we filed a motion for partial stay of the consolidated action in favor of the Louisiana Municipal Police Employees Retirement System action. In April 2011, the

court granted the motions to dismiss the consolidated complaint with leave to amend.

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We are involved in various other lawsuits and claims arising in the ordinary course of business. These other matters are, in the opinion of management, immaterial both individually and in the aggregate with respect to our consolidated financial position, liquidity or results of operations. Because of the uncertainties related to the incurrence, amount and range of loss on any pending litigation, investigation, inquiry or claim, management is currently unable to predict the ultimate outcome of any litigation, investigation, inquiry or claim, determine whether a liability has been incurred or make an estimate of the reasonably possible liability that could result from an unfavorable outcome. We believe however, that the liability, if any, resulting from the aggregate amount of uninsured damages for any outstanding litigation, investigation or claim will not have a material adverse effect on our consolidated financial position, liquidity or results of operations. However, an adverse ruling in a patent infringement lawsuit involving us could materially affect our ability to sell one or more of our products or could result in additional competition. In view of the unpredictable nature of such matters, we cannot provide any assurances regarding the outcome of any litigation, investigation, inquiry or claim to which we are a party or the impact on us of an adverse ruling in such matters.

Item 1A. ***Risk Factors***

The risk factors presented below update, and should be considered in addition to, the risk factors previously disclosed by us in Part I, Item 1A Risk Factors of our Annual Report on Form 10-K for the fiscal year ended December 31, 2010.

We may experience losses due to product liability claims, product recalls or corrections.

The design, development, manufacture and sale of our products involve an inherent risk of product liability or other claims by consumers and other third parties. We have in the past been, and continue to be, subject to various product liability claims and lawsuits. In addition, we have in the past and may in the future recall or issue field corrections related to our products due to manufacturing deficiencies, labeling errors or other safety or regulatory reasons. We cannot assure you that we will not in the future experience material losses due to product liability claims, lawsuits, product recalls or corrections.

As part of the Inamed acquisition, we assumed Inamed s product liability risks, including any product liability for its past and present manufacturing of breast implant products. The manufacture and sale of breast implant products has been and continues to be the subject of a significant number of product liability claims due to allegations that the medical devices cause disease or result in complications, rare lymphomas and other health conditions due to rupture, deflation or other product failure. Historically, other breast implant manufacturers that suffered such claims in the 1990 s were forced to cease operations or even to declare bankruptcy.

Additionally, FDA marketing approval for our silicone breast implants requires that:

- we monitor patients in our core study out to 10 years even if there has been explantation of the core device without replacement;
- patients in the core study receive magnetic resonance imaging tests, or MRIs, at seven and nine years;
- we conduct a large, 10-year post-approval study;
- we monitor patients in our adjunct study through the patients 5-year evaluation; and
- we conduct additional smaller evaluations, including a focus group aimed at ensuring patients are adequately informed about the risks of our silicone breast implants and that the format and content of patient labeling is adequate.

We are seeking marketing approval for other silicone breast implants in the United States, and if we obtain this approval, it may similarly be subject to significant restrictions and requirements, including the need for a patient registry, follow up MRIs and substantial post-market clinical trial commitments.

We also face a substantial risk of product liability claims from our eye care, neuromodulator, urology, skin care, obesity intervention and facial aesthetics products. Additionally, our pharmaceutical and medical device products may cause, or may appear to cause, serious adverse side

effects or potentially dangerous drug interactions if misused, improperly prescribed, improperly implanted or based on faulty surgical technique. We are subject to adverse event reporting regulations that require us to report to the FDA or similar bodies in other countries if our products are associated with a death or serious injury, even if there is no available evidence of a causal relationship between the adverse event and the product. Such reports may be publicly released by the FDA and other authorities. For instance, the FDA maintains a public database, known as the Manufacturer and User Facility Device Experience, or MAUDE, that posts reports of adverse events involving medical devices. The submission of an adverse event report for a pharmaceutical or medical device product to the FDA and its public release on MAUDE, or other public database, does not, by regulation, reflect a conclusion by us or the FDA that the product

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caused or contributed to the adverse event. However, as part of our post-market pharmacovigilance program, we routinely monitor the adverse event reports we receive to identify potential safety issues, known as signals, that may require us to take action with respect to the product, such as a recall or other market action, and/or amending our labeling to add the adverse reaction and/or a new warning or contraindication. The FDA and other regulatory authorities also monitor adverse event reports to identify safety signals, and may take action in connection with that monitoring, including the imposition on us of additional regulatory controls, such as the performance of costly post-approval clinical studies or revisions to our approved labeling, which could limit the indications or patient population for our products or could even lead to the withdrawal of a product from the market. We cannot assure you that the FDA will agree with our assessments of whether a safety signal exists for one of our products. Furthermore, any adverse publicity associated with adverse events for our products, and related post-market actions, could cause consumers to seek alternatives to our products, which may cause our sales to decline, even if our products are ultimately determined not to have been the primary cause of the adverse event.

Natural disasters and geo-political events could adversely affect our business.

We are a global company with sales and marketing subsidiaries in over 35 countries and are present in over 100 countries, as supplemented by distributors. The occurrence of one or more natural disasters, such as earthquakes, tsunamis, hurricanes, floods and tornados, or severe changes in geo-political events, such as wars, civil unrest or terrorist attacks in a country in which we operate or in which our suppliers or distributors are located could adversely affect our business and financial performance. Such events could result in physical damage to, or the complete loss of, properties or assets that are important to us or to our suppliers or distributors, changes in consumers income or purchasing patterns, temporary or long-term disruption in the supply of products to us, or disruption in the distribution of our products. Any such events and their consequences are unpredictable and could disrupt our operations or the operations of our suppliers or distributors and could have a significant and adverse effect on our business and results of operations.

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

The following table discloses the purchases of our equity securities during the first fiscal quarter of 2011.

Period	Total Number of Shares Purchased(1)	Average Price Paid per Share	Total Number of Shares Purchased as Part of Publicly Announced Plans or Programs	Maximum Number (or Approximate Dollar Value) of Shares that May Yet be Purchased Under the Plans or Programs(2)
January 1, 2011 to January 31, 2011	720,700	$ 70.48	720,700	15,759,002
February 1, 2011 to February 28, 2011	684,900	72.18	684,900	16,719,791
March 1, 2011 to March 31, 2011	883,500	70.85	883,500	15,908,401
Total	2,289,100	$ 71.14	2,289,100	N/A

(1) We maintain an evergreen stock repurchase program, which we first announced on September 28, 1993. Under the stock repurchase program, we may maintain up to 18.4 million repurchased shares in our treasury account at any one time. At March 31, 2011, we held approximately 2.5 million treasury shares under this program. Effective January 1, 2011, our current Rule 10b5-1 plan authorizes our broker to purchase our common stock traded in the open market pursuant to our evergreen stock repurchase program. The terms of the plan set forth a maximum limit of 4.0 million shares to be repurchased, certain quarterly maximum and minimum volume limits, and the plan is cancellable at any time in our sole discretion and in accordance with applicable insider trading laws.

(2) The share numbers reflect the maximum number of shares that may be purchased under our stock repurchase program and are as of the end of each of the respective periods.

Item 3. ***Defaults Upon Senior Securities***

None.

Item 4. ***(Removed and Reserved)***

Item 5. ***Other Information***

None.

Item 6. ***Exhibits***

Reference is made to the Exhibit Index included herein.

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SIGNATURE

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.

Date: May 5, 2011

ALLERGAN, INC.

/s/ Jeffrey L. Edwards
Jeffrey L. Edwards

Executive Vice President,

Finance and Business Development,

Chief Financial Officer

(Principal Financial Officer)

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ALLERGAN, INC.

EXHIBIT INDEX

Exhibit No.	Description
3.1	Amended and Restated Certificate of Incorporation of Allergan, Inc., as filed with the State of Delaware on May 4, 2011
3.2	Allergan, Inc. Amended and Restated Bylaws (incorporated by reference to Exhibit 3.1 to Allergan, Inc. s Current Report on Form 8-K filed on October 7, 2008)
10.1	Allergan, Inc. Pension Plan (Restated 2011) (incorporated by reference to Exhibit 10.20 to Allergan, Inc. s Annual Report on Form 10-K for the Fiscal Year ended December 31, 2010)
10.2	Allergan, Inc. Executive Severance Pay Plan (Effective January 2011) (incorporated by reference to Exhibit 10.1 to Allergan, Inc. s Current Report on Form 8-K filed on December 21, 2010)
10.3	Allergan, Inc. 2011 Executive Bonus Plan (incorporated by reference to Annex A to Allergan, Inc. s Proxy Statement filed on March 8, 2011)
10.4	Allergan, Inc. 2011 Management Bonus Plan (incorporated by reference to Exhibit 10.24 to Allergan, Inc. s Annual Report on Form 10-K for the Fiscal Year ended December 31, 2010)
10.5	Allergan, Inc. 2011 Incentive Award Plan (incorporated by reference to Annex B to Allergan, Inc. s Proxy Statement filed on March 8, 2011)
10.6	Form of Non-Qualified Stock Option Grant Notice for Employees under the Allergan, Inc. 2011 Incentive Award Plan
10.7	Form of Restricted Stock Award Grant Notice for Employees under the Allergan, Inc. 2011 Incentive Award Plan
10.8	Form of Restricted Stock Award Grant Notice for Employees (Management Bonus Plan) under the Allergan, Inc. 2011 Incentive Award Plan
10.9	Form of Restricted Stock Unit Award Grant Notice for Employees under the Allergan, Inc. 2011 Incentive Award Plan
10.10	Form of Restricted Stock Unit Award Grant Notice for Employees (Management Bonus Plan) under the Allergan, Inc. 2011 Incentive Award Plan
10.11	Form of Restricted Stock Unit Award Grant Notice for Non-Employee Directors under the Allergan, Inc. 2011 Incentive Award Plan
10.12	Collaboration Agreement, dated as of January 28, 2011, among MAP Pharmaceuticals, Inc., Allergan USA, Inc., Allergan Sales, LLC and Allergan, Inc.* (incorporated by reference to Exhibit 10.55 to Allergan, Inc. s Annual Report on Form 10-K for the Fiscal Year ended December 31, 2010)
10.13	Co-Promotion Agreement, dated as of January 28, 2011, among MAP Pharmaceuticals, Inc., Allergan USA, Inc. and Allergan, Inc.* (incorporated by reference to Exhibit 10.56 to Allergan, Inc. s Annual Report on Form 10-K for the Fiscal Year ended December 31, 2010)
31.1	Certification of Principal Executive Officer Required Under Rule 13a-14(a) of the Securities Exchange Act of 1934, as amended
31.2	Certification of Principal Financial Officer Required Under Rule 13a-14(a) of the Securities Exchange Act of 1934, as amended
32	Certification of Principal Executive Officer and Principal Financial Officer Required Under Rule 13a-14(b) of the Securities Exchange Act of 1934, as amended, and 18 U.S.C. Section 1350

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Exhibit No.	Description
101	The following financial statements are from Allergan, Inc. s Report on Form 10-Q for the Quarter ended March 31, 2011, formatted in XBRL (eXtensible Business Reporting Language): (i) Unaudited Condensed Consolidated Statements of Earnings, (ii) Unaudited Condensed Consolidated Balance Sheets; (iii) Unaudited Condensed Consolidated Statements of Cash Flows; and (iv) Notes to Unaudited Condensed Consolidated Financial Statements

* Confidential treatment was requested with respect to the omitted portions of this Exhibit, which portions have been filed separately with the Securities and Exchange Commission and which portions were granted confidential treatment