Thousands of dollars, except per share amounts and numbers of shares

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Summary of Significant Accounting Policies

Principles of Consolidation

The consolidated financial statements include the accounts of Becton, Dickinson and Company and its majority-owned subsidiaries (the "Company") after the elimination of intercompany transactions. The Company has no material interests in variable interest entities.

Cash Equivalents

Cash equivalents consist of all highly liquid investments with a maturity of three months or less at time of purchase.

Short-Term Investments

Short-term investments consist of time deposits with maturities greater than three months and less than one year when purchased.

Inventories

Inventories are stated at the lower of first-in, first-out cost or market.

Property, Plant and Equipment

Property, plant and equipment are stated at cost, less accumulated depreciation and amortization. Depreciation and amortization are principally provided on the straight-line basis over estimated useful lives, which range from 20 to 45 years for buildings, four to 10 years for machinery and equipment and two to 17 years for leasehold improvements. Depreciation and amortization expense was \$305,510, \$280,357 and \$262,956 in fiscal 2008, 2007 and 2006, respectively.

Goodwill and Other Intangible Assets

Goodwill is reviewed annually for impairment in accordance with Statement of Financial Accounting Standards ("SFAS") No. 142, "Goodwill and Other Intangible Assets." In reviewing goodwill for impairment, potential impairment is identified by comparing the fair value of a reporting unit, estimated using an income approach, with its carrying value. Core and developed technology is amortized over periods ranging from 15 to 20 years, using the straight-line method. Both goodwill and core and developed technology arise from acquisitions. Other intangibles with finite useful lives, which include patents, are amortized over periods principally ranging from two to 40 years, using the straight-line method. These intangibles, including core and developed technology,

are periodically reviewed when impairment indicators are present to assess recoverability from future operations using undiscounted cash flows in accordance with SFAS No. 144, "Accounting for the Impairment or Disposal of Long-Lived Assets." To the extent carrying value exceeds the undiscounted cash flows, an impairment loss is recognized in operating results based upon the excess of the carrying value over fair value. Other intangibles also include certain trademarks that are considered to have indefinite lives, as they are expected to generate cash flows indefinitely, and are reviewed annually for impairment.

Capitalized Software

Capitalized software, including costs for software developed or obtained for internal use is stated at cost, less accumulated amortization. Amortization expense is principally provided on the straight-line basis over estimated useful lives, which do not exceed 10 years. Amortization expense was \$56,652, \$66,386 and \$66,037 for 2008, 2007 and 2006, respectively.

Foreign Currency Translation

Generally, the net assets of foreign operations are translated into U.S. dollars using current exchange rates. The U.S. dollar results that arise from such translation, as well as exchange gains and losses on intercompany balances of a long-term investment nature, are included in the foreign currency translation adjustments in Accumulated other comprehensive (loss) income.

Revenue Recognition

Revenue from product sales is recognized when title and risk of loss pass to the customer. The Company recognizes revenue for certain instruments sold from the Biosciences segment upon installation at a customer's site, as installation of these instruments is considered a significant post-delivery obligation. For certain sales arrangements, primarily in the U.S., with multiple deliverables, revenue and cost of products sold are recognized at the completion of each deliverable: shipment, installation and training, in accordance with Emerging Issues Task Force Issue No. 00-21, "Revenue Arrangements with Multiple Deliverables." These sales agreements are divided into separate units of accounting. Revenue is recognized upon the completion of each deliverable based on the relative fair values of items delivered. Fair values are generally determined based on sales of the individual deliverables to other third parties.

The Company's domestic businesses sell products primarily to distributors who resell the products to end-user customers. Rebates are provided to distributors that sell to end-user customers at prices determined under a contract between the Company and the end-user customer. Provisions for rebates, as well as sales discounts and returns, are accounted for as a reduction of revenues when revenue is recognized.

Shipping and Handling Costs

Shipping and handling costs are included in Selling and administrative expense. Shipping expense was \$276,370, \$243,263 and \$219,788 in 2008, 2007 and 2006, respectively.

Derivative Financial Instruments

In accordance with SFAS No. 133, "Accounting for Derivative Instruments and Hedging Activities," as amended, all derivatives are recorded in the balance sheet at fair value and changes in fair value are recognized currently in earnings unless specific hedge accounting criteria are met.

Derivative financial instruments are utilized by the Company in the management of its foreign currency and interest rate exposures. The Company hedges its foreign currency exposures by entering into offsetting forward exchange contracts and currency options when it deems appropriate. The Company utilizes interest rate swaps and forward rate agreements to manage its exposure to fluctuating interest rates. The Company does not use derivative financial instruments for trading or speculative purposes.

Any deferred gains or losses associated with derivative instruments, which on infrequent occasions may be terminated prior to maturity, are recognized in income in the period in which the underlying hedged transaction is recognized. In the event a designated hedged item is sold, extinguished or matures prior to the termination of the related derivative instrument, such instrument would be closed and the resultant gain or loss would be recognized in income.

Income Taxes

United States income taxes are not provided on undistributed earnings of foreign subsidiaries where such undistributed earnings are indefinitely reinvested outside the United States. Deferred taxes are provided for earnings of foreign subsidiaries when those earnings are not considered indefinitely reinvested. Income taxes are provided and tax credits are recognized based on tax laws enacted at the dates of the financial statements.

The Company maintains valuation allowances where it is more likely than not that all or a portion of a deferred tax asset will not be realized. Changes in valuation allowances are included in our tax provision in the period of change. In determining whether a valuation allowance is warranted, management evaluates factors such as prior earnings history, expected future earnings, carry back and carry forward periods and tax strategies that could potentially enhance the likelihood of the realization of a deferred tax asset.

Earnings per Share

Basic earnings per share are computed based on the weighted average number of common shares outstanding. Diluted earnings per share reflect the potential dilution that could occur if securities or other contracts to issue common stock were exercised or converted into common stock.

Use of Estimates

The preparation of financial statements in conformity with U.S. generally accepted accounting principles requires management to make estimates and assumptions. These estimates or assumptions affect reported assets, liabilities, revenues and expenses as reflected in the consolidated financial statements. Actual results could differ from these estimates.

Share-Based Compensation

The Company accounts for all share-based compensation under SFAS No. 123 (revised 2004) – "Share-Based Payment" ("SFAS No. 123(R)"). This statement requires the recognition of the fair value of share-based compensation in net income. Compensation expense is recognized on a straight-line basis over the requisite service period, which is generally the vesting period.

2 Accounting Changes

On October 1, 2007, the Company adopted Financial Accounting Standards Board Interpretation No. 48 "Accounting for Uncertainty in Income Taxes, an interpretation of FASB Statement No. 109" ("FIN 48"). FIN 48 provides guidance for the financial statement recognition and measurement of a tax position taken or expected to be taken in a tax return. As a result of the implementation of FIN 48, the Company recognized a \$5,083 increase in its existing liability for uncertain tax positions, with a corresponding decrease to the October 1, 2007 retained earnings balance. The Company also reclassified the total amount of unrecognized tax benefits of \$71,782 from a current liability account (Accrued expenses) to a non-current liability account (Deferred Income Taxes and Other) on the Consolidated Balance Sheets, in accordance with FIN 48 as of October 1, 2007. If the Company were to recognize the unrecognized tax benefits, the effective tax rate would be favorably impacted. The Company does not anticipate any significant changes over the next 12 months to the amount of unrecognized tax benefits.

The Company includes interest and penalties associated with unrecognized tax benefits as a component of the Income tax provision on the Consolidated Statements of Income. As of October 1, 2007, accrued interest and penalties related to unrecognized tax benefits, included in the total amount, were \$9,388.

The Company conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of tax jurisdictions. The U.S. Internal Revenue Service ("IRS") has completed its audit for the tax years through 2002; however, the tax years 2000 through 2002 remain open, with a single issue being considered in the IRS administrative appeals process. For the Company's other major tax jurisdictions where it conducts business, the Company's tax years are generally open after 2002.

In September 2006, the Financial Accounting Standards Board (the "FASB") issued SFAS No. 158, "Employers' Accounting for Defined Benefit Pension and Other Postretirement Plans, an amendment of FASB Statements No. 87, 88, 106 and 132(R)" ("SFAS No. 158"). SFAS No. 158 requires the Company to recognize the overfunded or underfunded status of a defined benefit postretirement plan as an asset or liability in its Consolidated Balance Sheet and to recognize changes in the funded status in the year in which the changes occur through comprehensive income.

SFAS No. 158 also requires the funded status of a plan to be measured as of the balance sheet date and provides for additional disclosure requirements. The Company adopted SFAS No. 158 on September 30, 2007. SFAS No. 158 did not change the measurement date of the Company's plans as the plans are measured at its fiscal year-end. See Note 5 regarding the Company's adoption of SFAS No. 158.

In March 2005, the FASB issued Interpretation No. 47 "Accounting for Conditional Asset Retirement Obligations" ("FIN 47"). FIN 47 clarifies that the term "conditional asset retirement obligation" as used in SFAS No. 143, "Accounting for Asset Retirement Obligations" refers to a legal obligation to perform an asset retirement activity in which the timing and/or method of settlement are conditional on a future event that may or may not be within the control of the Company. Accordingly, the Company is required to recognize a liability for the fair value of a conditional asset retirement obligation if the fair value can be reasonably estimated. The Company adopted this interpretation in the fourth quarter of 2006. The adoption of FIN 47 did not have a material impact on BD's consolidated financial statements.

Adoption of New Accounting Standard

In September 2006, the FASB issued SFAS No. 157, "Fair Value Measurements" ("SFAS No. 157"). SFAS No. 157 defines fair value, establishes a framework for measuring fair value in GAAP, and expands disclosures about fair value measurements. This Statement is effective for the Company beginning October 1, 2008, and applies to interim periods. The Company does not anticipate the implementation of this Statement will be material to the consolidated financial position or results of operations.

In February 2007, the FASB issued SFAS No. 159, "The Fair Value Option for Financial Assets and Financial Liabilities" ("SFAS No. 159"). SFAS No. 159 permits entities to choose to measure many financial instruments and certain other items at fair value. Under SFAS No. 159, the decision to measure items as fair value is made at specified election dates on an instrument-by-instrument basis and is irrevocable. Entities electing the fair value option are required to recognize changes in fair value in earnings and to expense upfront costs and fees associated with the item for which the fair value option is elected. This Statement would be effective for the Company, if adopted, beginning October 1, 2008; however, the Company does not anticipate adopting this Statement.

3 Acquisitions and Divestitures

Cytopeia

On May 12, 2008, the Company acquired 100% of the outstanding stock of Cytopeia, Inc., a privately-held corporation that develops and markets advanced flow cytometry cell sorting instruments. The acquisition advances the Company's position in rapidly emerging areas of cell-based research, such as cell therapy research, stem cell research, drug discovery and development, and marine biology. The acquisition was accounted for under the purchase method of accounting and the results of operations of Cytopeia were included in the Biosciences segment's results as of the acquisition date. Pro forma information was not provided as the acquisition did not have a material effect on the Company's consolidated results. The purchase price was \$42,914 in cash, including transaction costs. Cash assumed as of the valuation date was \$1,655. The purchase price was allocated based upon the fair values of the assets and liabilities acquired. The allocation is based upon the information available as of September 30, 2008 and may be adjusted should further information become available. Additional information that may become subsequently available includes, but is not limited to, changes in the value of deferred tax assets and liabilities. The purchase price allocation resulted in a deferred tax asset of \$4,290, core and developed technology of \$20,000, deferred tax liabilities of \$7,904, primarily associated with core and developed technology; and other net assets of \$3,713, primarily consisting of accounts receivable and inventory. Core and developed technology will be amortized on a straight-line basis over its estimated useful life of approximately 15 years. The excess of the purchase price over the fair value of the assets acquired of \$22,815 was recorded as goodwill. The primary item that generated goodwill is the value of the Company's access to new technologies and capabilities related to cell therapy research. No portion of this goodwill is expected to be deductible for tax purposes.

TriPath

On December 20, 2006, the Company acquired the outstanding shares (approximately 93.8%) of TriPath Imaging, Inc. ("TriPath") which it did not previously own. TriPath develops, manufactures, markets and sells innovative solutions to improve the clinical management of cancer, including detection, diagnosis, staging and treatment. The acquisition advances the Company's position in cancer diagnostics. The acquisition was accounted for under the purchase method of accounting and the results of operations of TriPath were included in the Company's results as of the acquisition date.

Pro forma information was not provided as the acquisition did not have a material effect on the Company's consolidated results. The purchase price was \$361,883 in cash, including transaction costs and other consideration. The purchase price was allocated based upon the fair values of the assets and liabilities acquired. The allocation of the purchase price resulted in deferred tax assets of \$75,261 primarily consisting of net operating loss carry-forwards and credits; core and developed technology of \$135,097; deferred tax liabilities of \$52,662 primarily associated with other intangible assets; and other net assets of \$56,736 consisting primarily of cash and trade receivables. Core and developed technology will be amortized on a straight-line basis over its estimated useful life of approximately 15 years. The excess of the purchase price over the fair value of the assets acquired of \$32,712 was recorded as goodwill. The primary items that generated goodwill are the value of expanded product opportunities in oncology that are aligned with and complement ongoing research programs at the Company. The goodwill was allocated to the Diagnostics segment and is not deductible for tax purposes. As a result of settling a preacquisition legal contingency in the fourth quarter of 2007, the Company received an upfront cash payment of \$7,167. The effects of this payment, as well as other minor purchase accounting adjustments, are reflected in the purchase price allocation detailed above.

In connection with the acquisition, the Company also incurred a non-deductible charge of \$114,739 for acquired in-process research and development. This charge, based on fair value, is associated with three projects: molecular Pap test, breast staging, and ovarian cancer tests. These projects had not yet reached technological feasibility and did not have alternative future use at the acquisition date. The portion of the charge allocated to each of these projects was \$75,992, \$18,764 and \$19,983, respectively.

The molecular Pap test uses proprietary molecular biomarkers and reagents that are intended to allow for the primary screening of cervical cancer. The addition of biomarkers is intended to improve sensitivity to allow the clinician to find disease more reliably. In February 2008, the Company ceased activities on the clinical trial for this product. The Company presently anticipates having a molecular Pap test commercially available both in the U.S. and outside the U.S. in fiscal year 2012, assuming successful completion of new clinical trials and attainment of approval from FDA.

The breast cancer project, using proprietary biomarkers and reagents, is intended to aid in disease discovery in its earliest stages. Tests developed in this program will also be run on the multiplex testing platform discussed below.

Since the acquisition, the Company changed the focus in this program from staging assay development to screening assay development. The Company anticipates having a breast assay both in the U.S. and outside the U.S. in fiscal year 2013, assuming successful completion of clinical trials and attainment of approval from FDA.

The ovarian cancer project, using proprietary biomarkers and reagents in a multiplex format, is intended to allow for earlier stage detection of cancer. Information the Company expects to obtain from tests developed in this project should allow clinicians to begin treatment sooner, which should lead to improved outcomes overall. In addition, the Company signed a development and supply agreement for a multiplex testing platform to allow for the simultaneous testing of multiple markers from a small volume of serum. The Company anticipates having an ovarian monitoring test commercially available in the U.S. in fiscal year 2010, assuming successful completion of clinical trials and attainment of approval from FDA. Screening tests are expected to be commercially available approximately two years thereafter.

The fair values of these projects were determined based upon the present value of projected cash flows utilizing an income approach reflecting the appropriate risk-adjusted discount rate based on the applicable technological and commercial risk of each project. These cash flows also took into account the income and expenses associated with the further development and commercialization of the underlying products. The range of discount rates assigned to the projects was 22 to 30 percent and gave consideration to the underlying risk relative to the developed technology, the overall commercial and technical risk, and the probabilities of success for each of the projects. The ongoing activity associated with each of these projects is not expected to be material to the Company's research and development expense.

Plasso

On May 4, 2007, the Company acquired all of the outstanding shares of Plasso Technology, Ltd. ("Plasso"), a privately-held company that is developing the next generation of surface-critical research tools utilizing functional coating technology for applications in glycomics and cell culture, for \$10,425 in cash including transaction costs. In connection with the acquisition, the Company incurred a non-deductible charge of \$7,394 for acquired in-process research and development associated with Plasso's technology, for which, at the acquisition date, technological feasibility had not been established and no alternative future use existed. Because Plasso was a development stage company that had not commenced its planned principal operations, the transaction was accounted for as an acquisition of assets rather than as a business combination and, therefore, goodwill was not recorded.

GeneOhm

On February 14, 2006, the Company acquired all the outstanding stock of GeneOhm Sciences, Inc. ("GeneOhm"), a company that develops molecular diagnostic testing for the rapid detection of bacterial organisms, including those known to cause healthcare-associated infections. The acquisition provides the Company with expanded entry into the emerging field of healthcare-associated infections. The acquisition was accounted for under the purchase method of accounting and the results of operations of GeneOhm were included in the Company's results as of the acquisition date. Pro forma information was not provided as the acquisition did not have a material effect on the Company's consolidated results. The purchase price consisted of an up-front cash payment of \$232,542, including transaction costs. The purchase price was allocated based upon the fair values of the assets and liabilities acquired. The allocation of the purchase price resulted in deferred tax assets of \$32,423 consisting of net operating loss carry forwards and credits; other intangible assets, primarily core and developed technology, of \$91,043; deferred tax liabilities of \$29,626 associated with other intangible assets, and other net assets of \$3,750. Core and developed technology will be amortized on a straight-line basis over its estimated useful life of approximately 15 years. The excess of the purchase price over the fair value of the assets acquired of \$81,652 was recorded as goodwill. The primary items that generated goodwill are the value of synergies in microbiology research and the expansion of product offerings in molecular diagnostics. The goodwill was allocated to the Diagnostics segment and is not deductible for tax purposes. In connection with the acquisition, the Company also incurred a non-deductible charge of \$53,300 for acquired in-process research and development. This charge, based on fair value, is associated with several products that have not reached technological feasibility and do not have alternative future use at the acquisition date. The fair value of each product was determined based upon the present value of projected cash flows utilizing an income approach reflecting the appropriate risk-adjusted discount rate based on the applicable technological and commercial risk of each product. These cash flows took into account the income and expenses associated with the further development and commercialization of the underlying products. The ongoing activity associated with each of these products is not material to the Company's research and development expense.

BGM

On September 28, 2006, the Company announced a plan to exit the blood glucose monitoring ("BGM") market. In accordance with the plan, distribution of the *BD Logic* Blood Glucose Monitor was immediately discontinued. BD continued to distribute test strips for its customers through December 2007. The decision to exit the BGM market was made following an evaluation of the future outlook for the product line. The Company recorded a pre-tax charge of \$63,414 in 2006 in connection with its decision to exit the BGM product line. This charge consisted of \$5,352 related to estimated customer sales returns, \$31,602 related to the write-off of inventory and related purchase commitments, \$14,052 related to long-lived asset write-downs, and \$12,408 related to severance and other exit costs.

During the first quarter of fiscal 2007, the Company received an unsolicited offer for the purchase of the BGM product line. On December 11, 2006, the Company sold the product line for \$19,971 and recognized a pre-tax gain on sale of \$15,226. During 2007, adjustments of \$9,319 were made to reduce sales returns and other accruals related to obligations that remained with the Company upon divestiture of the product line. Additionally, the Company received a payment of \$4,675, which represented the resolution of a contingency with a former supplier. Following the sale, the Company's prior period Consolidated Statements of Income and Cash Flows and related disclosures have been restated to separately present the results of the BGM product line as discontinued operations.

Other

In August 2005, the Company completed the sale of the Clontech unit of the Biosciences segment for \$62,100 and recognized a gain on sale of \$13,336 (\$28,533 after taxes). Clontech's results of operations were reported as discontinued operations in 2006 in the accompanying Consolidated Statements of Income and Cash Flows.

Results of discontinued operations for the years ended September 30 were as follows:

	2008	2007	2006
Revenues	\$2,587	\$33,086	\$ 96,811
(Loss) income from discontinued			
operations before income taxes	(1,489)	49,108	(95,653) ^(A)
Income tax benefit (provision)	567	(15,242)	32,823
(Loss) income from discontinued			
operations, net	\$ (922)	\$33,866	\$(62,830)(A)

⁽A) Includes post-closing charges of \$4,708 (\$3,311 after taxes) related to the divestiture of Clontech.

4 Other Intangible Assets

Other intangible assets at September 30 consisted of:

	2008		2007		
	Gross		Gross		
	Carrying	Accumulated	Carrying	Accumulated	
	Amount	Amortization	Amount	Amortization	
Amortized intangible					
assets					
Core and developed					
technology	\$548,974	\$200,443	\$548,995	\$174,216	
Patents, trademarks,					
and other	297,321	216,697	289,920	203,037	
	\$846,295	\$417,140	\$838,915	\$377,253	
Unamortized intangib	le				
assets					
Trademarks	\$ 9,051		\$ 9,055		

Intangible amortization expense was \$54,217, \$46,607 and \$34,843 in 2008, 2007 and 2006, respectively. The estimated aggregate amortization expense for the fiscal years ending September 30, 2009 to 2013 are as follows: 2009 – \$51,900; 2010 – \$50,000; 2011 – \$48,100; 2012 – \$45,200; 2013 – \$43,800.

5 Renefit Plans

The Company has defined benefit pension plans covering substantially all of its employees in the United States and certain foreign locations. The Company also provides certain postretirement healthcare and life insurance benefits to qualifying domestic retirees. Postretirement healthcare and life insurance benefit plans in foreign countries are not material. The measurement date used for the Company's employee benefit plans is September 30.

During 2007, the Company redesigned its U.S. pension plans to provide for a cash benefit formula by offering a one-time, irrevocable election to existing employees to change to this provision and mandating all new employees hired after April 1, 2007 to participate in the new formula. The Company also amended its other postretirement benefits plan to provide that new hires, as of April 1, 2007 or later, will no longer be eligible for company subsidized benefits. These amendments did not have a material impact on the net pension and postretirement cost of the Company in 2007.

Net pension and other postretirement cost for the years ended September 30 included the following components:

	Pension Plans		Othe	Other Postretirement Bene		
	2008	2007	2006	2008	2007	2006
Service cost	\$66,440	\$69,869	\$74,111	\$ 4,648	\$ 4,386	\$ 4,164
Interest cost	81,939	75,728	71,997	14,906	14,608	14,873
Expected return on plan assets	(97,218)	(88,527)	(80,063)	_	_	_
Amortization of prior service cost	(1,066)	348	309	(6,232)	(6,233)	(6,233)
Amortization of loss	8,256	17,507	27,932	3,962	5,795	7,127
Amortization of net obligation	(112)	(92)	(70)	_	_	_
Settlements	602	_	_	_	_	_
	\$58,841	\$74,833	\$94,216	\$17,284	\$18,556	\$19,931

Net pension cost attributable to foreign plans included in the preceding table was \$20,072, \$21,156 and \$18,639 in 2008, 2007 and 2006, respectively.

Effective September 30, 2007, the Company adopted the recognition and disclosure provisions of SFAS No. 158, which requires the Company to recognize on a prospective basis the funded status of its pension and other postretirement benefit plans in the Consolidated Balance Sheet with a corresponding adjustment to Accumulated other comprehensive (loss) income. The incremental effect of adopting SFAS No. 158 was a \$209,695 reduction in Shareholders' Equity, net of deferred taxes as of September 30, 2007.

The change in benefit obligation, change in fair value of plan assets, funded status and amounts recognized in the Consolidated Balance Sheets for these plans were as follows:

	Pension Plans		Other Postretirement Benefits	
	2008	2007	2008	2007
Change in benefit obligation:				
Beginning obligation	\$1,394,430	\$1,384,667	\$ 245,971	\$ 255,726
Service cost	66,440	69,869	4,648	4,386
Interest cost	81,939	75,728	14,906	14,608
Plan amendments	_	(16,586)	_	_
Benefits paid	(71,517)	(97,671)	(22,303)	(25,411)
Actuarial gain	(181,968)	(63,519)	(47,605)	(11,818)
Other, includes translation	(16,868)	41,942	5,629	8,480
Benefit obligation at September 30	\$1,272,456	\$1,394,430	\$ 201,246	\$ 245,971
Change in fair value of plan assets:				
Beginning fair value	\$1,296,169	\$1,124,565	s —	\$ —
Actual return on plan assets	(224,777)	138,446	_	_
Employer contribution	114,924	96,952	_	_
Benefits paid	(71,517)	(97,671)	_	_
Other, includes translation	(14,833)	33,877	_	_
Plan assets at September 30	\$1,099,966	\$1,296,169	s —	\$ —
Funded Status at September 30:				
Unfunded benefit obligation	\$ (172,490)	\$ (98,261)	\$(201,246)	\$(245,971)
Amounts recognized in the Consolidated				
Balance Sheets at September 30:				
Other	\$ 2,841	\$ 32,710	\$ —	\$ —
Salaries, wages and related items	(5,006)	(2,668)	(19,427)	(20,067)
Long-term Employee Benefit Obligations	(170,325)	(128,303)	(181,819)	(225,904)
Net amount recognized	\$ (172,490)	\$ (98,261)	\$(201,246)	\$(245,971)
Amounts recognized in Accumulated other comprehensive				
(loss) income before income taxes at September 30:				
Net transition obligation	\$ 951	\$ 1,156	\$ (243)	\$ —
Prior service credit	9,018	10,086	456	6,688
Net actuarial loss	(359,793)	(238,144)	(9,992)	(62,194)
Net amount recognized	\$ (349,824)	\$ (226,902)	\$ (9,779)	\$ (55,506)

Foreign pension plan assets at fair value included in the preceding table were \$303,146 and \$359,291 at September 30, 2008 and 2007, respectively. The foreign pension plan projected benefit obligations were \$417,344 and \$430,265 at September 30, 2008 and 2007, respectively.

The projected benefit obligation, accumulated benefit obligation and fair value of plan assets for the pension plans with accumulated benefit obligations in excess of plan assets were \$260,253, \$227,820 and \$135,442, respectively as of September 30, 2008, and \$96,723, \$76,398 and \$14,685, respectively as of September 30, 2007.

The estimated net actuarial loss and prior service credit for pension benefits that will be amortized from Accumulated other comprehensive (loss) income into net pension costs over the next fiscal year are expected to be \$17,905 and \$1,093, respectively. The estimated net actuarial gain and prior service credit for other postretirement benefits that will be amortized from Accumulated other comprehensive (loss) income into net other postretirement costs over the next fiscal year are expected to be \$131 and \$463, respectively.

The weighted average assumptions used in determining pension plan information were as follows:

	2008	2007	2006
Net Cost			
Discount rate:			
U.S. plans ^(A)	6.35%	5.95%	5.50%
Foreign plans	5.32	4.65	4.19
Expected return on plan assets:			
U.S. plans ^(A)	8.00	8.00	8.00
Foreign plans	6.42	6.42	6.02
Rate of compensation increase:			
U.S. plans ^(A)	4.50	4.50	4.25
Foreign plans	3.45	3.08	2.92
Benefit Obligation			
Discount rate:			
U.S. plans ^(A)	8.00	6.35	5.95
Foreign plans	5.98	5.32	4.65
Rate of compensation increase:			
U.S. plans(A)	4.50	4.50	4.50
Foreign plans	3.56	3.45	3.08

(A) Also used to determine other postretirement and postemployment benefit plan information.

At September 30, 2008 the assumed healthcare trend rates were 8% pre and post age 65, gradually decreasing to an ultimate rate of 5% beginning in 2015. At September 30, 2007 the corresponding assumed healthcare trend rates were 9% pre and post age 65, gradually decreasing to an ultimate rate of 5% beginning in 2012. A one percentage point increase in assumed healthcare cost trend rates in each year would increase the accumulated postretirement benefit obligation as of September 30, 2008 by \$10,943 and the aggregate of the service cost and interest cost components of 2008 annual expense by \$762. A one percentage point decrease in the assumed healthcare cost trend rates in each year would decrease the accumulated postretirement benefit obligation as of September 30, 2008 by \$10,073 and the aggregate of the 2008 service cost and interest cost by \$667.

Expected Funding

The Company's funding policy for its defined benefit pension plans is to contribute amounts sufficient to meet legal funding requirements, plus any additional amounts that may be appropriate considering the funded status of the plans, tax consequences, the cash flow generated by the Company and other factors. While the Company will not be required to fund any of its pension plans in 2009, the Company made a discretionary contribution to its U.S. pension plan in October 2008 of \$75,000.

Expected benefit payments are as follows:

	Pension Plans	Other Postretirement Benefits
2009	\$105,137	\$ 19,427
2010	77,257	19,714
2011	82,588	19,870
2012	89,884	20,019
2013	95,246	20,178
2014-2018	607,818	101,109

Expected receipts of the subsidy under the Medicare Prescription Drug Improvement and Modernization Act of 2003, which are not reflected in the expected other postretirement benefit payments included in the preceding table, are as follows: 2009, \$2,372; 2010, \$2,480; 2011, \$2,574; 2012, \$2,646; 2013, \$2,695; 2014-2018, \$13,314.

The Company's asset allocations for its defined benefit pension plans at September 30 were as follows:

	2008	2007
Equity securities	55.1%	64.5%
Debt securities	35.7	33.1
Other (primarily cash)	9.2	2.4
	100.0%	100.0%

Investment Strategy

The Company's investment objective is to achieve superior returns on plan assets, subject to a prudent level of portfolio risk, for the purpose of enhancing the security of benefits for participants. The Company's investments include a broad range of equity and fixed-income securities. These investments are diversified in terms of domestic and international equity securities, short-term and long-term securities, growth and value styles, as well as small and large capitalization stocks. The Company's target allocation percentages are as follows: equity securities (58% - 69%); fixed-income securities (31% - 39%); and cash (0% - 3%). Equity securities are held for their expected high return and excess return over inflation. Fixed-income securities are held for diversification relative to equities. The plans may also hold cash to meet liquidity requirements. Due to short-term fluctuations in market conditions, allocation percentages may temporarily deviate from these target allocation percentages before a rebalancing occurs. Investment risks and returns are measured and monitored on an on-going basis through annual liability measurements and quarterly investment portfolio reviews to determine whether the asset allocation targets continue to represent an appropriate balance of expected risk and reward. The Company is in the process of determining the optimal deployment of its discretionary contributions of \$75,000 made in both September and October 2008.

The expected rate of return on plan assets is based upon expectations of long-term average rates of return to be achieved by the underlying investment portfolios. In establishing this assumption, the Company considers many factors, including historical assumptions compared with actual results; benchmark data; expected returns on various plan asset classes, as well as current and expected asset allocations.

Postemployment Benefits

The Company utilizes a service-based approach in applying SFAS No. 112, "Employers' Accounting for Postemployment Benefits – an amendment of FASB Statements No. 5 and 43," for most of its postemployment benefits. This approach recognizes that actuarial gains and losses may result from experience that differs from baseline assumptions.

Postemployment benefit costs for the years ended September 30 included the following components:

	2008	2007	2006
Service cost	\$11,276	\$10,449	\$10,148
Interest cost	5,643	5,116	4,946
Amortization of prior service cost	159	1,654	1,654
Amortization of loss	6,686	6,895	8,548
	\$23,764	\$24,114	\$25,296

The unfunded status of the postemployment benefit plans, which are not funded, was \$76,286 and \$101,514 at September 30, 2008 and 2007, respectively. The amounts recognized in Accumulated other comprehensive (loss) income before income taxes for the net actuarial loss was \$26,014 and \$57,110 at September 30, 2008 and 2007, respectively. The estimated net actuarial loss that will be amortized from the Accumulated other comprehensive (loss) income into postemployment benefit cost over the next fiscal year is \$2,627.

Savings Incentive Plan

The Company has a voluntary defined contribution plan ("Savings Incentive Plan") covering eligible employees in the United States. In connection with the redesign of the U.S. pension and postretirement benefit plans, effective July 1, 2007, the Company amended its Savings Incentive Plan increasing the amount of the Company matching contribution for eligible employees to 75% of employees' contributions, up to a maximum of 4.5% of each employee's eligible compensation. Prior to that date, the Company matched 50% of employees' contributions, up to a maximum of 3% of each employee's salary. The cost of the Savings Incentive Plan was \$31,526 in 2008, \$21,878 in 2007 and \$16,626 in 2006. The Company guarantees employees' contributions to the fixed income fund of the Savings Incentive Plan, which consists of diversified money market instruments. The amount guaranteed was \$175,344 at September 30, 2008.

6 Income

The provision for income taxes from continuing operations for the years ended September 30 consisted of:

	2008	2007	2006
Current:			
Federal	\$269,638	\$307,072	\$281,784
State and local, including Puerto Rico	13,872	21,669	12,004
Foreign	150,009	134,526	125,289
	433,519	463,267	419,077
Deferred:			
Domestic	12,384	(94,306)	(101,651)
Foreign	(20,214)	(21,183)	(6,634)
	(7,830)	(115,489)	(108,285)
	\$425,689	\$347,778	\$310,792

The components of Income From Continuing Operations Before Income Taxes for the years ended September 30 consisted of:

	2008	2007	2006
Domestic, including Puerto Rico	\$ 790,894	\$ 550,750	\$ 466,655
Foreign	762,713	653,195	659,247
	\$1,553,607	\$1,203,945	\$1,125,902

Deferred tax assets and liabilities are netted on the balance sheet by separate tax jurisdictions. At September 30, 2008 and 2007, net current deferred tax assets of \$211,188 and \$168,305, respectively, were included in Prepaid expenses, deferred taxes and other. Net non-current deferred tax assets of \$85,311 and \$168,251, respectively, were included in Other. Net current deferred tax liabilities of \$2,985 and \$6,136, respectively, were included in Current Liabilities -Income taxes. Net non-current deferred tax liabilities of \$35,519 and \$37,121, respectively, were included in Deferred Income Taxes and Other. Deferred taxes are not provided on undistributed earnings of foreign subsidiaries that are indefinitely reinvested. At September 30, 2008, the cumulative amount of such undistributed earnings indefinitely reinvested outside the United States was \$2.1 billion. Determining the tax liability that would arise if these earnings were remitted is not practicable. Deferred taxes are provided for earnings outside the United States when those earnings are not considered indefinitely reinvested.

On October 1, 2007, BD adopted the provisions of FIN 48, "Accounting for Uncertainty in Income Taxes." Under this Interpretation, in order to recognize an uncertain tax benefit, the taxpayer must be more likely than not of sustaining the position, and the measurement of the benefit is calculated as the largest amount that is more than 50 percent likely to be realized upon resolution of the benefit. Adoption of this Interpretation did not have a material impact on BD's financial position. The following table summarizes the gross amounts of unrecognized tax benefits without regard to reduction in tax liabilities or additions to deferred tax assets and liabilities if such unrecognized tax benefits were settled:

October 1, 2007	\$71,782
Increase due to current year tax positions	5,411
Increase due to prior year tax positions	535
Decrease due to settlements and lapse of statute of limitations	(8,030)
September 30, 2008	\$69,698

The total amount of unrecognized tax benefits, if recognized, would favorably impact the effective tax rate. Included in the above total is approximately \$10,946 of interest and penalties, of which approximately \$1,558 are reflected in the current year statement of operations. BD does not expect significant changes in the aggregate amount of unrecognized tax benefits that may occur within the next twelve months, other than tax settlements.

The Company conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of tax jurisdictions. The IRS has completed its audit for the tax years through 2002; however, the tax years 2000 through 2002 remain open, with a single issue being considered in the IRS administrative appeals process. For the Company's other major tax jurisdictions where it conducts business, the Company's tax years are generally open after 2002.

In October 2004, the American Jobs Creations Act of 2004 (the "AJCA") was signed into law. The AJCA created a temporary incentive for U.S. multinationals to repatriate accumulated income earned outside the United States. As a result of the passage of the AJCA, the Company revisited its policy of indefinite reinvestment of foreign earnings and made a decision to repatriate approximately \$1.3 billion in 2006 pursuant to its approved repatriation plan. During 2006, the Company repatriated approximately \$1.3 billion in accordance with its planned repatriation under the AJCA. The actual tax charge associated with this repatriation was \$65,768.

Deferred income taxes at September 30 consisted of:

		2008		2007
	Assets	Liabilities	Assets	Liabilities
Compensation and benefits	\$297,933	\$ —	\$301,118	\$ —
Property and equipment	_	206,503	_	190,979
Loss and credit carryforwards	175,341	_	193,981	_
Other	281,279	189,741	172,740	83,538
	754,553	396,244	667,839	274,517
Valuation allowance	(100,314)	_	(100,023)	_
	\$654,239	\$396,244	\$567,816	\$274,517

Valuation allowances have been established for capital loss carryforwards, state deferred tax assets, net of federal tax, related to net operating losses and credits and other deferred tax assets for which the Company has determined it is more likely than not that these benefits will not be realized. At September 30, 2008, the Company had deferred state tax assets for net state operating losses and credit carryforwards of \$49,965 for which a valuation allowance of \$33,860 has been established due to the uncertainty of generating sufficient taxable income in the state jurisdictions to utilize the deferred tax assets before they principally expire between 2009 and 2014. In 2007, a previously established valuation allowance of approximately \$19,700 related to state tax credit carryforwards was reversed and included in the state and local income tax line item in the following rate reconciliation table. The Company also has federal and state capital loss carryforward deferred tax assets of \$51,428 for which a full valuation allowance has been established due to the uncertainty of recognizing the benefit from these losses before they principally expire in 2010.

A reconciliation of the federal statutory tax rate to the Company's effective tax rate was as follows:

	2008	2007	2006
Federal statutory tax rate	35.0%	35.0%	35.0%
State and local income taxes,			
net of federal tax benefit	1.4	.2	.6
Effect of foreign and			
Puerto Rico earnings and			
foreign tax credits	(8.1)	(9.2)	(7.4)
Effect of Research, Domestic Production			
Activities, Extraterritorial Income			
tax benefits	(8.0)	(0.5)	(1.3)
Acquired in-process research and			
development	_	3.6	1.8
Repatriation of foreign earnings under			
the AJCA	_	_	(1.1)
Other, net	(0.1)	(0.2)	_
	27.4%	28.9%	27.6%

The approximate dollar and diluted earnings per share amounts of tax reductions related to tax holidays in various countries in which the Company does business were: 2008 - \$84,600 and \$0.33; 2007 - \$80,300 and \$0.32; and 2006 - \$70,000 and \$0.27. The tax holidays expire at various dates through 2023.

The Company made income tax payments, net of refunds, of \$330,709 in 2008, \$345,049 in 2007 and \$398,808 in 2006.

Supplemental Financial Information

Other (Expense) Income, Net

Other (expense) income, net in 2008 was \$(1,484), which primarily included foreign exchange losses (inclusive of hedging costs) of \$(10,303), partially offset by equity investment income of \$4,642 and income from license and other agreements of \$3,386.

Other (expense) income, net in 2007 was \$944, which primarily included income from license and other agreements of \$6,128, partially offset by net write downs of certain investments of \$(5,538) and foreign exchange losses (inclusive of hedging costs) of \$(4,191).

Other (expense) income, net in 2006 was \$(8,762), which primarily included net write downs of certain investments of \$(11,046) and foreign exchange losses (inclusive of hedging costs) of \$(5,142), partially offset by income from license and other agreements of \$4,281.

Trade Receivables, Net

Allowances for doubtful accounts and cash discounts netted against trade receivables were \$35,614 and \$39,650 at September 30, 2008 and 2007, respectively.

Inventories

Inventories at September 30 consisted of:

	2008	2007
Materials	\$ 162,726	\$142,484
Work in process	203,926	195,155
Finished products	713,774	714,320
	\$1,080,426	\$1,051,959

Property, Plant and Equipment, Net

Property, Plant and Equipment, Net at September 30 consisted of:

	2008	2007
Land	\$ 93,339	\$ 79,368
Buildings	1,803,620	1,597,356
Machinery, equipment and fixtures	3,822,785	3,596,781
Leasehold improvements	78,251	80,610
	5,797,995	5,354,115
Less accumulated depreciation and amortization	3,053,521	2,856,777
	\$2,744,474	\$2,497,338



Short-term debt at September 30 consisted of:

	2008	2007
Loans Payable		
Domestic	\$200,000	\$200,000
Foreign	992	6,768
Current portion of long-term debt	320	866
	\$201,312	\$207,634

Domestic loans payable consist of commercial paper. Foreign loans payable consist of short-term borrowings from financial institutions. The weighted average interest rates for Short-term debt were 2.3% and 5.2% at September 30, 2008 and 2007, respectively. During 2008, we amended our \$1 billion syndicated credit facility to extend its expiration date from December 2011 to December 2012. This credit facility provides backup support for the commercial paper program and can also be used for other general corporate purposes. It includes a restrictive covenant that requires a minimum interest coverage ratio, with which the Company was in compliance at September 30, 2008. There were no borrowings outstanding under the facility at September 30, 2008. In addition, the Company had short-term foreign lines of credit pursuant to informal arrangements of approximately \$145,000 at September 30, 2008, almost all of which was unused.

Long-Term Debt at September 30 consisted of:

	2008	2007
Domestic notes due through 2013 (average year-end		
interest rate: 2.4% - 2008; 4.3% - 2007)	\$ 8,130	\$ 9,801
7.15% Notes due October 1, 2009	205,372	205,914
4.55% Notes due April 15, 2013	198,940	198,734
4.90% Notes due April 15, 2018	205,734	206,214
7.00% Debentures due August 1, 2027	168,000	168,000
6.70% Debentures due August 1, 2028	167,050	167,050
	\$953,226	\$955,713

Long-term debt balances as of September 30, 2008 and 2007 have been impacted by certain interest rate swaps that have been designated as fair value hedges, as discussed in Note 9.

The aggregate annual maturities of long-term debt during the fiscal years ending September 30, 2010 to 2013 are as follows: 2010 - \$205,457; 2011 - \$22; 2012 - \$20; 2013 - \$206,943.

The Company capitalizes interest costs as a component of the cost of construction in progress. A summary of interest costs for the years ended September 30 were as follows:

	2008	2007	2006
Charged to operations	\$36,343	\$46,420	\$66,046
Capitalized	29,862	27,528	19,955
	\$66,205	\$73,948	\$86,001

Interest paid, net of amounts capitalized, was \$36,222 in 2008, \$50,730 in 2007 and \$62,514 in 2006.

9 Financial Instruments

Foreign Exchange Derivatives

The Company uses foreign exchange forward contracts and currency options to reduce the effect of fluctuating foreign exchange rates on certain foreign currency denominated receivables and payables and third party product sales. Gains and losses on the derivatives are intended to offset gains and losses on the hedged transaction. The Company's foreign currency risk exposure is in Europe, Asia-Pacific, Canada, Japan, and Latin America.

The Company hedges substantially all of its transactional foreign exchange exposures, primarily intercompany payables and receivables, through the use of forward contracts and currency options with maturities of less than 12 months. Gains or losses on these contracts are largely offset by gains and losses on the underlying hedged items. These foreign exchange contracts do not qualify for hedge accounting.

In addition, the Company enters into forward and option contracts to hedge certain forecasted sales that are denominated in foreign currencies. These contracts are designated as cash flow hedges and are effective as hedges of these revenues. These contracts are intended to reduce the risk that the Company's cash flows from certain third party transactions will be adversely affected by changes in foreign currency exchange rates. Changes in the effective portion of the fair value of these contracts are included in other comprehensive income until the hedged sales transactions are recognized in earnings. Once the hedged transaction occurs, the gain or loss on the contract is recognized from Accumulated other comprehensive (loss) income to revenues. The Company recorded hedge net gains, exclusive of hedging costs, of \$1,177, \$6,911 and \$8,242 to revenues in 2008, 2007 and 2006, respectively. Revenues in 2008, 2007 and 2006 are net of hedging costs of \$12,037, \$15,136 and \$12,508, respectively, related to the purchased option contracts. The Company records in Other income (expense), net, the premium of the forward contracts, which is excluded from the assessment of hedge effectiveness. The net premium was \$562 in 2006. All outstanding contracts that were designated as cash flow hedges as of September 30, 2008 will mature by September 30, 2009. At September 30, 2008 and 2007, Accumulated other comprehensive (loss) income included unrealized gains of \$37,786 and unrealized losses of \$4,994, respectively, net of tax, relating to foreign exchange derivatives that have been designated as cash flow hedges.

Interest Rate Derivatives

The Company's policy is to manage interest cost using a mix of fixed and floating rate debt. The Company has entered into interest rate swaps in which it agrees to exchange, at specified intervals, the difference between fixed and floating interest amounts calculated by reference to an agreed-upon notional principal amount. These swaps are designated as either fair value or cash flow hedges. For fair value hedges, changes in the fair value of the interest rate swaps offset changes in the fair value of the fixed rate debt due to changes in market interest rates. For cash flow hedges, changes in the fair value of the interest rate swaps are offset by amounts recorded in other comprehensive (loss) income. There was no ineffective portion to the hedges recognized in earnings during the period. If interest rate derivatives designated as cash flow hedges mature or are terminated, then the balance in other comprehensive (loss) income attributable to those derivatives is reclassified into earnings over the remaining life of the hedged debt. The amount that will be reclassified and recorded in Interest expense within the next 12 months is \$1,763.

At September 30, 2008 and 2007, Accumulated other comprehensive (loss) income included an unrealized loss of \$10,306 and \$11,397, respectively, net of tax, relating to interest rate derivatives that have been designated as cash flow hedges.

Fair Value of Financial Instruments

Cash equivalents, short-term investments and short-term debt are carried at cost, which approximates fair value. Equity securities, where a readily determinable market value exists, are classified as available-for-sale securities. Available-for-sale securities are carried at fair value, with unrecognized gains and losses reported in other comprehensive (loss) income, net of taxes. Losses on available-for-sale securities are recognized when a loss is determined to be other than temporary or when realized.

The fair value of forward exchange contracts and currency options were estimated based on market prices, where available, or dealer quotes. The fair value of certain long-term debt is based on redemption value. The estimated fair values of the Company's financial instruments at September 30 were as follows:

	20	800	2007	
	Carrying	Fair	Carrying	Fair
	Value	Value	Value	Value
Assets:				
Currency options(A)	s —	s —	\$3,982	\$3,982
Forward exchange contracts(A)	78,337	78,337	8,007	8,007
Interest rate swaps ^(A)	5,372	5,372	5,914	5,914
Equity securities	239	239	1,291	1,291
Liabilities:				
Forward exchange contracts(B)	29,647	29,647	8,968	8,968
Long-term debt	953,226	907,293	955,713	949,490

(A) Included in Prepaid expenses, deferred taxes and other.

(B) Included in Accrued Expenses.

Concentration of Credit Risk

Cash deposits in excess of amounts covered by government-provided insurance are exposed to loss in the event of non-performance by financial institutions. The Company does maintain cash deposits in excess of government-provided insurance limits. Substantially all of the Company's trade receivables are due from public and private entities involved in the healthcare industry. Due to the large size and diversity of the Company's customer base, concentrations of credit risk with respect to trade receivables are limited. The Company does not normally require collateral. The Company is exposed to credit loss in the event of nonperformance by financial institutions with which it conducts business. However, this loss is limited to the amounts, if any, by which the obliga-

tions of the counterparty to the financial instrument contract exceed the obligations of the Company. The Company also minimizes exposure to credit risk by dealing with a diversified group of major financial institutions.

10 Shareholders' Equity

Changes in certain components of shareholders' equity were as follows:

	Common					
	Stock	Capital in				
	Issued at	Excess of	Retained	Deferred	Treasu	ry Stock
	Par Value	Par Value	Earnings	Compensation	Shares	Amount
Balance at September 30, 2005	\$332,662	\$615,846	\$4,805,852	\$10,280	(84,977,933)	\$(2,297,493)
Net income			752,280			
Cash dividends:						
Common (\$.86 per share)			(212,435)			
Common stock issued for:						
Share-based compensation plans, net		148,342			5,066,384	49,057
Business acquisitions		734			15,864	156
Share-based compensation		108,613				
Common stock held in trusts, net				854	(17,275)	(854)
Repurchase of common stock					(7,281,100)	(448,882)
Balance at September 30, 2006	\$332,662	\$873,535	\$5,345,697	\$11,134	(87,194,060)	\$(2,698,016)
Net income			890,033			
Cash dividends:						
Common (\$.98 per share)			(239,943)			
Common stock issued for:						
Share-based compensation plans, net		143,420			4,380,724	43,213
Business acquisitions		707			10,812	105
Share-based compensation		107,706				
Common stock held in trusts, net				1,071	(70,542)	(1,071)
Repurchase of common stock					(5,952,000)	(450,124)
Balance at September 30, 2007	\$332,662	\$1,125,368	\$5,995,787	\$12,205	(88,825,066)	\$(3,105,893)
Net income			1,126,996			
Cash dividends:						
Common (\$1.14 per share)			(279,110)			
Common stock issued for:						
Share-based compensation plans, net		132,372			4,649,160	25,866
Business acquisitions		1,206			16,327	118
Share-based compensation		100,585				
Common stock held in trusts, net				2,489	(169,307)	(2,489)
Repurchase of common stock					(5,255,900)	(450,000)
Cumulative effect for adoption of FIN 48			(5,084)			
Balance at September 30, 2008	\$332,662	\$1,359,531	\$6,838,589	\$14,694	(89,584,786)	\$(3,532,398)

Common stock held in trusts represents rabbi trusts in connection with deferred compensation under the Company's employee salary and bonus deferral plan and directors' deferral plan.

1 1 Accumulated Other Comprehensive (Loss) Income

The components of Accumulated other comprehensive (loss) income were as follows:

	2008	2007
Foreign currency translation adjustments	\$157,089	\$237,394
Benefit plans adjustment	(261,457)	(218,595)
Unrealized loss on investments	(622)	(580)
Unrealized gains (losses) on cash flow hedges	27,480	(16,391)
	\$ (77,510)	\$ 1,828

The income tax benefit recorded in fiscal years 2008 and 2007 for the unrealized gains on investments was \$25 and \$6,524, respectively. The income tax provision (benefit) recorded in fiscal years 2008 and 2007 for cash flow hedges was \$26,889 and \$(1,247), respectively. The income tax benefit recorded in fiscal year 2008 for defined benefit pension and postretirement plans was \$3,439. The income tax provision recorded in fiscal year 2007 for the minimum pension liability adjustment was \$2,050. Income taxes are generally not provided for translation adjustments.

The unrealized losses on cash flow hedges included in other comprehensive (loss) income for 2008 and 2007 are net of reclassification adjustments of \$6,733 and \$5,099, net of tax, respectively, for realized net hedge losses recorded to revenues. These amounts had been included in Accumulated other comprehensive (loss) income in prior periods. The tax benefits associated with these reclassification adjustments in 2008 and 2007 were \$4,127 and \$3,126, respectively.

12 Commitments and Contingencies

Commitments

Rental expense for all operating leases amounted to \$70,300 in 2008, \$68,100 in 2007, and \$63,400 in 2006. Future minimum rental commitments on noncancelable leases are as follows: 2009 - \$48,600; 2010 - \$37,900; 2011 - \$30,300; 2012 - \$24,500; 2013 - \$21,100 and an aggregate of \$27,200 thereafter.

As of September 30, 2008, the Company has certain future purchase commitments aggregating to approximately \$505,000, which will be expended over the next several years.

Contingencies

The Company is named as a defendant in five purported class action suits brought on behalf of direct purchasers of the Company's products, such as distributors, alleging that the Company violated federal antitrust laws, resulting in the charging of higher prices for the Company's products to the plaintiff and other purported class members. The cases filed are as follows: Louisiana Wholesale Drug Company, Inc., et. al. vs. Becton Dickinson and Company (Civil Action No. 05-1602, U.S. District Court, Newark, New Jersey), filed on March 25, 2005; SAJ Distributors, Inc. et. al. vs. Becton Dickinson & Co. (Case 2:05-CV-04763-JD, U.S. District Court, Eastern District of Pennsylvania), filed on September 6, 2005; Dik Drug Company, et. al. vs. Becton, Dickinson and Company (Case No. 2:05-CV-04465, U.S. District Court, Newark, New Jersey), filed on September 12, 2005; American Sales Company, Inc. et. al. vs. Becton, Dickinson & Co. (Case No. 2:05-CV-05212-CRM, U.S. District Court, Eastern District of Pennsylvania), filed on October 3, 2005; and Park Surgical Co. Inc. et. al. vs. Becton, Dickinson and Company (Case 2:05-CV-05678-CMR, U.S. District Court, Eastern District of Pennsylvania), filed on October 26, 2005.

The actions brought by Louisiana Wholesale Drug Company and Dik Drug Company in New Jersey have been consolidated under the caption "In re Hypodermic Products Antitrust Litigation."

The Company is also named as a defendant in four purported class action suits brought on behalf of indirect purchasers of the Company's products, alleging that the Company violated federal antitrust laws, resulting in the charging of higher prices for the Company's products to the plaintiff and other purported class members. The cases filed are as follows: Jabo's Pharmacy, Inc., et. al. v. Becton Dickinson & Company (Case No. 2:05-CV-00162, U.S. District Court, Greenville, Tennessee), filed on June 7, 2005; Drug Mart Tallman, Inc., et. al. v. Becton Dickinson and Company (Case No. 2:06-CV-00174, U.S. District Court, Newark, New Jersey), filed on January 17, 2006; Medstar v. Becton Dickinson (Case No. 06-CV-03258-JLL (RJH), U.S. District Court, Newark, New Jersey), filed on May 18, 2006; and The Hebrew Home for the Aged at Riverdale v. Becton Dickinson and Company (Case No. 07-CV-2544, U.S. District Court, Southern District of New York), filed on March 28, 2007. A fifth purported class action on behalf of indirect purchasers International Multiple Sclerosis Management Practice v. Becton Dickinson & Company (Case No. 2:07-cv-10602, U.S. District Court, Newark, New Jersey), filed on April 5, 2007) was voluntarily withdrawn by the plaintiff.

The plaintiffs in each of the antitrust class action lawsuits seek monetary damages. All of the antitrust class action lawsuits have been consolidated for pre-trial purposes in a Multi-District Litigation (MDL) in federal court in New Jersey.

On June 6, 2006, UltiMed, Inc., a Minnesota company, filed suit against the Company in the U.S. District Court in Minneapolis, Minnesota (*UltiMed, Inc. v. Becton, Dickinson and Company* (06CV2266)). The plaintiff alleges, among other things, that the Company excluded the plaintiff from the market for home use insulin syringes by entering into anticompetitive contracts in violation of federal and state antitrust laws. The plaintiff seeks money damages and injunctive relief.

In June 2007, Retractable Technologies, Inc. ("RTI") filed a complaint against the Company under the caption *Retractable Technologies*, *Inc. vs. Becton Dickinson and Company* (Civil Action No. 2:07-cv-250, U.S. District Court, Eastern District of Texas). RTI alleges that the *BD Integra* syringes infringe patents licensed exclusively to RTI. In its complaint, RTI also alleges that the Company engaged in false advertising with respect to certain of the Company's safety-engineered products in violation of the Lanham Act;

acted to exclude RTI from various product markets and to maintain its market share through, among other things, exclusionary contracts in violation of state and federal antitrust laws; and engaged in unfair competition. In January 2008, the court granted the Company's motion to sever the patent and non-patent claims into separate cases. The non-patent claims have been stayed, pending resolution of RTI's patent claims. The trial on the patent claims is currently scheduled to commence in March 2009. RTI seeks money damages and injunctive relief. On April 1, 2008, RTI filed a complaint against BD under the caption Retractable Technologies, Inc. and Thomas J. Shaw v. Becton Dickinson and Company (Civil Action No. 2:08-cv-141, U.S. District Court, Eastern District of Texas). RTI alleges that the BD Integra syringes infringe another patent licensed exclusively to RTI. RTI seeks money damages and injunctive relief. On August 29, 2008, the court ordered the consolidation of these two cases.

The Company, along with another manufacturer and several medical product distributors, is named as a defendant in two product liability lawsuits relating to healthcare workers who allegedly sustained accidental needlesticks, but have not become infected with any disease. Generally, these actions allege that healthcare workers have sustained needlesticks using hollow-bore needle devices manufactured by the Company and, as a result, require medical testing, counseling and/or treatment. In some cases, these actions additionally allege that the healthcare workers have sustained mental anguish. Plaintiffs seek money damages in all of these actions. The Company had previously been named as a defendant in nine similar suits relating to healthcare workers who allegedly sustained accidental needlesticks, each of which has either been dismissed with prejudice or voluntarily withdrawn. Regarding the two pending suits:

- In Ohio, *Grant vs. Becton Dickinson et al.* (Case No. 98CVB075616, Franklin County Court), on September 21, 2006, the Ohio Court of Appeals reversed the trial court's grant of class certification. The matter has been remanded to the trial court for a determination of whether the class can be redefined.
- In South Carolina, a suit has been filed on behalf of an unspecified number of healthcare workers seeking class action certification in state court under the caption *Bales vs. Becton Dickinson et. al.* (Case No. 98-CP-40-4343, Richland County Court of Common Pleas), filed on November 25, 1998.

The Company continues to oppose class action certification in the pending cases, including pursuing all appropriate rights of appeal.

The Company, along with a number of other manufacturers, was named as a defendant in approximately 524 product liability lawsuits in various state and Federal courts related to natural rubber latex gloves which the Company ceased manufacturing in 1995. Cases pending in Federal court are being coordinated under the matter In re Latex Gloves Products Liability Litigation (MDL Docket No. 1148) in Philadelphia, and analogous procedures have been implemented in the state courts of California, Pennsylvania, New Jersey and New York. Generally, these actions allege that medical personnel have suffered allergic reactions ranging from skin irritation to anaphylaxis as a result of exposure to medical gloves containing natural rubber latex. Since the inception of this litigation, 467 of these cases have been closed with no liability to the Company, and 46 cases have been settled for an aggregate de minimis amount.

On May 28, 2004, Therasense, Inc. ("Therasense") filed suit against the Company (Therasense, Inc. and Abbott Laboratories v. Nova Biomedical Corporation and Becton, Dickinson and Company (Case Number: C 04-02123 WDA, U.S. District Court, Northern District of California)) asserting that the Company's blood glucose monitoring products infringe four Therasense patents and seeking money damages. On August 10, 2004, in response to a motion filed by Therasense in the U.S. District Court for the District of Massachusetts, the court transferred to the court in California an action previously filed by the Company against Therasense requesting a declaratory judgment that the Company's products do not infringe the Therasense patents and that the Therasense patents are invalid. On April 4, 2008, the Court granted the Company summary judgment with respect to two of the patents asserted against the Company, finding no infringement by the Company. On June 24, 2008, the Court ruled that a third patent asserted against the Company was invalid and unenforceable. On August 8, 2008, a jury delivered a verdict in the Company's favor, finding that the last of the four patents asserted against the Company was invalid. Abbott/Therasense have appealed some of these decisions, and it is possible that other decisions will also be appealed after the Court rules on post-trial motions.

On September 19, 2007, the Company was served with a qui tam complaint filed by a private party against the Company in the United States District Court, Northern District of Texas, alleging violations of the Federal False Claims Act ("FCA") and the Texas False Claims Act (the "TFCA") (U.S. ex rel Fitzgerald v. BD et al. (Civil Action No. 3:03-CV-1589, U.S. District Court, Northern District of Texas). The suit alleges that a group purchasing organization's practices with its suppliers, including the Company, inflated the costs of healthcare reimbursement. Under the FCA, the United States Department of Justice, Civil Division has a certain period of time in which to decide whether to join the claim against the Company as an additional plaintiff; if not, the private plaintiff is free to pursue the claim on its own. A similar process is followed under the TFCA. To the Company's knowledge, no decision has yet been made by the Civil Division or the State of Texas whether to join this claim. In September 2008, the Court dismissed certain of the plaintiff's claims, but denied the Company's motion to dismiss with respect to other claims.

The Company believes that it has meritorious defenses to each of the above-mentioned suits pending against the Company and is engaged in a vigorous defense of each of these matters.

The Company is also involved both as a plaintiff and a defendant in other legal proceedings and claims that arise in the ordinary course of business.

The Company is a party to a number of Federal proceedings in the United States brought under the Comprehensive Environment Response, Compensation and Liability Act, also known as "Superfund," and similar state laws. The affected sites are in varying stages of development. In some instances, the remedy has been completed, while in others, environmental studies are commencing. For all sites, there are other potentially responsible parties that may be jointly or severally liable to pay all cleanup costs.

Given the uncertain nature of litigation generally, the Company is not able in all cases to estimate the amount or range of loss that could result from an unfavorable outcome of the litigation to which the Company is a party. In accordance with U.S. generally accepted accounting principles, the Company establishes accruals to the extent probable future losses are estimable (in the case of environmental matters, without considering possible third-party recoveries). In view of the uncertainties discussed above, the Company could incur charges in excess of any currently established accruals and, to the extent available, excess liability insurance. In the opinion of management, any such future charges, individually or in the aggregate, could have a material adverse effect on the Company's consolidated results of operations and consolidated cash flows.

13

Share-Based Compensation

The Company grants share-based awards under the 2004 Employee and Director Equity-Based Compensation Plan ("2004 Plan"), which provides long-term incentive compensation to employees and directors consisting of: stock appreciation rights ("SARs"), stock options, performance-based restricted stock units, time-vested restricted stock units and other stock awards. In 2008, 2007 and 2006, the compensation expense for these plans charged to income was \$100,585, \$107,706 and \$108,613, respectively, and the associated income tax benefit recognized was \$36,236, \$37,179 and \$35,155, respectively.

Stock Appreciation Rights

SARs represent the right to receive, upon exercise, shares of common stock having a value equal to the difference between the market price of common stock on the date of exercise and the exercise price on the date of grant. SARs vest over a four-year period and have a ten-year term, similar to the previously granted stock options. The fair value was estimated on the date of grant using a lattice-based binomial option valuation model that uses the following weighted-average assumptions in 2008 and 2007: risk-free interest rate of 3.83% and 4.56%, respectively; expected volatility of 27% and 28%, respectively; expected dividend yield of 1.35% and 1.37%, respectively, and expected life of 6.5 years for both years. Expected volatility is based upon historical volatility for the Company's common stock and other factors. The expected term of SARs granted is derived from the output of the model, using assumed exercise rates based on historical exercise and termination patterns, and represents the period of time that SARs granted are expected to be outstanding. The risk-free interest rate used is based upon the published U.S. Treasury yield curve in effect at the time of grant for instruments with a similar life. The dividend yield is based upon the most recently declared quarterly dividend as of the grant date. The weighted average grant date fair value of SARs granted during 2008 and 2007 was \$24.92 and \$22.66, respectively. The total intrinsic value of SARs exercised during 2008 was \$2,122. The Company issued 17,873 shares during 2008 to satisfy the SARs exercised.

A summary of SARs outstanding as of September 30, 2008, and changes during the year then ended is as follows:

			Weighted	
		Weighted	Average	
		Average	Remaining	Aggregate
		Exercise	Contractual	Intrinsic
	SARs	Price	Term (Years)	Value
Balance at October 1	3,164,729	\$65.26		
Granted	1,445,508	84.33		
Exercised	(88,681)	62.33		
Forfeited, canceled				
or expired	(178,380)	71.11		
Balance at September 30	4,343,176	\$71.43	8.12	\$44,048
Vested and expected to				
vest at September 30	4,026,657	\$71.20	8.10	\$41,613
Exercisable at	·			
September 30	1,177,988	\$63.57	7.49	\$19,702

Stock options

All stock option grants are for a ten-year term. Stock options issued after November 2001 vest over a four-year period. Stock options issued prior to November 2001 vested over a three-year period. Stock options granted in 2005 were valued based on the grant date fair value of those awards, using a lattice-based binomial option valuation model that used the following weighted-average assumptions: risk-free interest rate of 3.93%; expected volatility of 29%; expected dividend yield of 1.28% and expected life of 6.5 years.

A summary of stock options outstanding as of September 30, 2008, and changes during the year then ended is as follows:

			Weighted
		Weighted	Average
		Average	Remaining Aggregate
	Stock	Exercise	Contractual Intrinsic
	Options	Price	Term (Years) Value
Balance at October 1	13,997,747	\$35.81	
Granted	_	_	
Exercised	(3,643,415)	33.75	
Forfeited, canceled			
or expired	(100,343)	39.09	
Balance at			
September 30	10,253,989	\$36.51	4.07 \$448,611
Vested and expected			
to vest at			
September 30	10,218,280	\$36.45	4.06 \$447,689
Exercisable at			
September 30	9,896,895	\$35.86	4.00 \$439,395

Cash received from the exercising of stock options in 2008, 2007 and 2006 was \$122,977, \$134,133 and \$147,831, respectively. The actual tax benefit realized for tax deductions from stock option exercises totaled \$62,230, \$59,491 and \$48,751, respectively. The total intrinsic value of stock options exercised during the years 2008, 2007 and 2006 was \$191,627, \$187,537 and \$168,752, respectively.

Performance-Based Restricted Stock Units

Performance-based restricted stock units cliff vest three years after the date of grant. These units are tied to the Company's performance against pre-established targets, including its average growth rate of consolidated revenues and average return on invested capital, over a three-year performance period. Under the Company's long-term incentive program, the actual payout under these awards may vary from zero to 250% of an employee's target payout, based on the Company's actual performance over the three-year performance period. The fair value is based on the market price of the Company's stock on the date of grant. Compensation cost initially recognized assumes that the target payout level will be achieved and is adjusted for subsequent changes in the expected outcome of performance-related conditions.

A summary of performance-based restricted stock units outstanding as of September 30, 2008, and changes during the year then ended is as follows:

		Weighted
		Average
	Stock	Grant Date
	Units	Fair Value
Balance at October 1	3,883,955	\$60.23
Granted	891,622	84.33
Vested	(671,208)	53.70
Forfeited or canceled	(937,074)	54.91
Balance at September 30 ^(A)	3,167,295	\$69.98
Expected to vest at September 30 ^(B)	1,338,925	\$69.24

- (A) Based on 170% to 250% of the target payout, depending on year of grant.
- (B) Net of expected forfeited units and units in excess of the expected performance payout of 194.157 and 1.634.213, respectively.

The weighted average grant date fair value of performance-based restricted stock units granted during the years 2007 and 2006 was \$71.72 and \$59.16, respectively. At September 30, 2008, the weighted average remaining contractual term of performance-based restricted stock units is 1.08 years.

Time-Vested Restricted Stock Units

Time-vested restricted stock units generally cliff vest three years after the date of grant, except for certain key executives of the Company, including the executive officers, for which such units generally vest one year following the employee's retirement. The related share-based compensation expense is recorded over the requisite service period, which is the vesting period or in the case of certain key executives is based on retirement eligibility. The fair value of all time-vested restricted stock units is based on the market value of the Company's stock on the date of grant.

A summary of time-vested restricted stock units outstanding as of September 30, 2008, and changes during the year then ended is as follows:

		Weighted
		Average
	Stock	Grant Date
	Units	Fair Value
Balance at October 1	1,618,082	\$61.11
Granted	469,625	84.42
Vested	(332,192)	55.72
Forfeited or canceled	(185,186)	60.00
Balance at September 30	1,570,329	\$69.35
Expected to vest at September 30	1,413,296	\$69.35

The weighted average grant date fair value of time-vested restricted stock units granted during the years 2007 and 2006 was \$72.20 and \$59.62, respectively. At September 30, 2008, the weighted average remaining contractual term of the time-vested restricted stock units is 1.90 years.

The amount of unrecognized compensation expense for all non-vested share-based awards as of September 30, 2008, is approximately \$106,872, which is expected to be recognized over a weighted-average remaining life of approximately 1.99 years. At September 30, 2008, 3,954,723 shares were authorized for future grants under the 2004 Plan.

The Company has a policy of satisfying share-based payments through either open market purchases or shares held in treasury. At September 30, 2008, the Company has sufficient shares held in treasury to satisfy these payments in 2009.

Other Stock Plans

The Company has a Stock Award Plan, which allows for grants of common shares to certain key employees. Distribution of 25% or more of each award is deferred until after retirement or involuntary termination, upon which the deferred portion of the award is distributable in five equal annual installments. The balance of the award is distributable over five years from the grant date, subject to certain conditions. In February 2004, this plan was terminated with respect to future grants upon the adoption of the 2004 Plan. At September 30, 2008 and 2007, awards for 161,145 and 214,206 shares, respectively, were outstanding.

The Company has a Restricted Stock Plan for Non-Employee Directors which reserves for issuance of 300,000 shares of the Company's common stock. No restricted shares were issued in 2008.

The Company has a Directors' Deferral Plan, which provides a means to defer director compensation, from time to time, on a deferred stock or cash basis. As of September 30, 2008, 97,881 shares were held in trust, of which 5,092 shares represented Directors' compensation in 2008, in accordance with the provisions of the plan. Under this plan, which is unfunded, directors have an unsecured contractual commitment from the Company.

The Company also has a Deferred Compensation Plan that allows certain highly-compensated employees, including executive officers, to defer salary, annual incentive awards and certain equity-based compensation. As of September 30, 2008, 454,316 shares were issuable under this plan.

14 Earnings per Share

The weighted average common shares used in the computations of basic and diluted earnings per share (shares in thousands) for the years ended September 30 were as follows:

	2008	2007	2006
Average common shares outstanding	244,323	244,929	247,067
Dilutive share equivalents from			
share-based plans	8,358	9,881	9,487
Average common and common			
equivalent shares outstanding -			
assuming dilution	252,681	254,810	256,554

15 Segment Data

The Company's organizational structure is based upon its three principal business segments: BD Medical ("Medical"), BD Diagnostics ("Diagnostics") and BD Biosciences ("Biosciences").

The principal product lines in the Medical segment include needles, syringes and intravenous catheters for medication delivery; safety-engineered and auto-disable devices; prefilled IV flush syringes; syringes and pen needles for the self-injection of insulin and other drugs used in the treatment of diabetes; prefillable drug delivery devices provided to pharmaceutical companies and sold to end-users as drug/device combinations; surgical blades/scalpels and regional anesthesia needles and trays; critical care monitoring devices; ophthalmic surgical instruments; sharps disposal containers; and home healthcare products. The principal products and services in the Diagnostics segment include integrated systems for specimen collection; an extensive line of safety-engineered specimen blood collection products and systems; plated media; automated blood culturing systems; molecular testing systems for sexually transmitted diseases and healthcare-associated infections; microorganism identification and drug susceptibility systems; liquid-based cytology systems for cervical cancer screening; and rapid diagnostic assays. The principal product lines in the Biosciences segment include fluorescence activated cell sorters and analyzers; cell imaging systems; monoclonal antibodies and kits for performing cell analysis; reagent systems for life sciences research; tools to aid in drug discovery and growth of tissue and cells; cell culture media supplements for biopharmaceutical manufacturing; and diagnostic assays.

The Company evaluates performance of its business segments based upon operating income. Segment operating income represents revenues reduced by product costs and operating expenses.

Distribution of products is primarily through independent distribution channels and directly to end-users by BD and independent sales representatives. Sales to a distributor that supplies products from the Medical and Diagnostics segments accounted for approximately 9% of revenues in 2008 and 2007. Sales to this distributor accounted for 11% of revenues in 2006. No other customer accounted for 10% or more of revenues in any of the three years presented.

Revenues ^(A)	2008	2007	2006
Medical	\$3,801,003	\$3,420,670	\$3,106,646
Diagnostics	2,159,811	1,905,105	1,715,090
Biosciences	1,195,096	1,033,933	916,281
	\$7,155,910	\$6,359,708	\$5,738,017
Segment Operating Income			
Medical	\$1,068,143	\$ 971,990	\$864,180
Diagnostics	525,747	342,778 ^(B)	390,355 ^(B)
Biosciences	333,662	258,806 ^(B)	221,925
Total Segment Operating Income	1,927,552	1,573,574	1,476,460
Unallocated Expenses(C)	(373,945)	(369,629)	(350,558)
Income From Continuing Operations			
Before Income Taxes	\$1,553,607	\$1,203,945	\$1,125,902
Segment Assets			
Medical	\$3,432,113	\$3,289,490	\$2,835,613
Diagnostics	1,887,261	1,843,654	1,485,959
Biosciences	933,105	817,000	727,634
Total Segment Assets	6,252,479	5,950,144	5,049,206
Corporate and All Other(D)	1,660,464	1,379,221	1,775,319
	\$7,912,943	\$7,329,365	\$6,824,525
Capital Expenditures			
Medical	\$378,786	\$ 352,696	\$ 268,669
Diagnostics	123,915	113,691	104,815
Biosciences	82,880	73,502	38,952
Corporate and All Other	16,400	16,505	44,631
	\$ 601,981	\$ 556,394	\$ 457,067
Depreciation and Amortization			
Medical	\$ 240,442	\$ 223,430	\$ 210,044
Diagnostics	150,202	138,936	116,072
Biosciences	75,809	68,889	63,383
Corporate and All Other	10,969	10,086	12,833
Corporate and Am Other			

⁽A) Intersegment revenues are not material.

Revenues by Organizational Units	2008	2007	2006
BD Medical			
Medical Surgical Systems	\$2,004,854	\$1,864,080	\$1,748,743
Diabetes Care	775,320	695,981	656,533
Pharmaceutical Systems	942,136	791,900	639,694
Ophthalmic Systems	78,693	68,709	61,676
	\$3,801,003	\$3,420,670	\$3,106,646
BD Diagnostics			
Preanalytical Systems	\$1,123,528	\$1,006,692	\$927,759
Diagnostic Systems	1,036,283	898,413	787,331
	\$2,159,811	\$1,905,105	\$1,715,090
BD Biosciences			
Cell Analysis(A)	\$ 900,511	\$ 756,031	\$ 660,196
Discovery Labware	294,585	277,902	256,085
	\$1,195,096	\$1,033,933	\$ 916,281
	\$7,155,910	\$6,359,708	\$5,738,017

(A) Cell Analysis consists of the Immunocytometry Systems and the Pharmingen organizational units that were previously reported separately.

Geographic Information

The countries in which the Company has local revenuegenerating operations have been combined into the following geographic areas: the United States (including Puerto Rico), Europe, and Other, which is composed of Canada, Latin America, Japan and Asia-Pacific.

Revenues to unaffiliated customers are based upon the source of the product shipment. Long-lived assets, which include net property, plant and equipment, are based upon physical location.

	2008	2007	2006
Revenues			
United States	\$3,184,806	\$3,033,005	\$2,739,344
Europe	2,488,956	2,047,388	1,762,782
Other	1,482,148	1,279,315	1,235,891
	\$7,155,910	\$6,359,708	\$5,738,017
Long-Lived Assets			
United States	\$2,179,544	\$2,172,327	\$1,934,994
Europe	1,135,379	1,106,284	893,495
Other	721,355	646,188	540,925
Corporate	261,990	274,000	269,858
	\$4,298,268	\$4,198,799	\$3,639,272

⁽B) Includes the acquired in-process research and development charges in 2007 related to the TriPath and Plasso acquisitions, and in 2006 related to the GeneOhm acquisition, as discussed in Note 3.

⁽C) Includes primarily interest, net; foreign exchange; corporate expenses and share-based compensation expense.

⁽D) Includes cash and investments and corporate assets.