BOSTON SCIENTIFIC CORP
Form 10-Q
August 03, 2016
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UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549
FORM 10-Q

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

For the quarterly period ended June 30, 2016
OR

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

Commission File No. 1-11083
BOSTON SCIENTIFIC CORPORATION
(Exact name of registrant as specified in its charter)

DELAWARE	04-2695240
(State or other jurisdiction of incorporation or organization)	(I.R.S. Employer Identification No.)

300 BOSTON SCIENTIFIC WAY, MARLBOROUGH, MASSACHUSETTS 01752-1234
(Address of principal executive offices) (zip code)
(508) 683-4000
(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 þ No o
Indicate by check mark whether the registrant has submitted electronically and posted on its corporate website, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§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 þ No o
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 þ	Accelerated filer o	Non-Accelerated filer o (Do not check if a smaller reporting company)	Smaller reporting company o

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes o No þ
Indicate the number of shares outstanding of each of the issuer's classes of common stock, as of the latest practicable date.

Class	Shares outstanding as of July 29, 2016
Common Stock, $.01 par value	1,360,743,339

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PART I
FINANCIAL INFORMATION

ITEM 1. CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

BOSTON SCIENTIFIC CORPORATION AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS (UNAUDITED)

	Three Months Ended June 30,		Six Months Ended June 30,	
in millions, except per share data	2016	2015	2016	2015
Net sales	$2,126	$1,843	$4,090	$3,611
Cost of products sold	639	540	1,211	1,060
Gross profit	1,487	1,303	2,879	2,551
Operating expenses:				
Selling, general and administrative expenses	779	700	1,497	1,367
Research and development expenses	222	220	431	412
Royalty expense	20	18	39	36
Amortization expense	135	116	271	229
Intangible asset impairment charges	—	9	—	9
Contingent consideration expense (benefit)	33	19	37	46
Restructuring charges	14	3	17	9
Litigation-related charges (credits)	618	(1)	628	192
Pension termination charges	—	—	—	8
	1,821	1,084	2,920	2,308
Operating income (loss)	(334)	219	(41)	243
Other income (expense):				
Interest expense	(59)	(106)	(118)	(167)
Other, net	(4)	(8)	(10)	(22)
Income (loss) before income taxes	(397)	105	(169)	54
Income tax expense (benefit)	(190)	3	(164)	(47)
Net income (loss)	$(207)	$102	$(5)	$101
Net income (loss) per common share — basic	$(0.15)	$0.08	$(0.00)	$0.08
Net income (loss) per common share — assuming dilution	$(0.15)	$0.08	$(0.00)	$0.07
Weighted-average shares outstanding				
Basic	1,357.4	1,341.3	1,353.9	1,337.5
Assuming dilution	1,357.4	1,361.8	1,353.9	1,359.7

See notes to the unaudited condensed consolidated financial statements.

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BOSTON SCIENTIFIC CORPORATION AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME (LOSS) (UNAUDITED)

	Three Months Ended June 30,		Six Months Ended June 30,	
(in millions)	2016	2015	2016	2015
Net income (loss)	$(207)	$102	$(5)	$101
Other comprehensive income (loss):				
Foreign currency translation adjustment	(21)	5	(5)	(30)
Net change in unrealized gains and losses on derivative financial instruments, net of tax	(84)	(43)	(153)	(15)
Net change in certain retirement plans, net of tax	—	—	—	5
Total other comprehensive income (loss)	(105)	(38)	(158)	(40)
Total comprehensive income (loss)	$(312)	$64	$(163)	$61

See notes to the unaudited condensed consolidated financial statements.

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BOSTON SCIENTIFIC CORPORATION AND SUBSIDIARIES
CONDENSED CONSOLIDATED BALANCE SHEETS

in millions, except share and per share data	As of June 30, 2016 (Unaudited)	December 31, 2015
ASSETS		
Current assets:		
Cash and cash equivalents	$438	$319
Trade accounts receivable, net	1,387	1,275
Inventories	981	1,016
Deferred and prepaid income taxes	78	496
Other current assets	446	365
Total current assets	3,330	3,471
Property, plant and equipment, net	1,487	1,490
Goodwill	6,475	6,473
Other intangible assets, net	5,930	6,194
Other long-term assets	616	505
TOTAL ASSETS	$17,838	$18,133
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Current debt obligations	$254	$3
Accounts payable	284	209
Accrued expenses	2,236	1,970
Other current liabilities	408	248
Total current liabilities	3,182	2,430
Long-term debt	5,173	5,674
Deferred income taxes	24	735
Other long-term liabilities	3,239	2,974
Commitments and contingencies		
Stockholders' equity		
Preferred stock, $.01 par value - authorized 50,000,000 shares, none issued and outstanding		
Common stock, $.01 par value - authorized 2,000,000,000 shares - issued 1,606,264,833 shares as of June 30, 2016 and 1,594,213,786 shares as of December 31, 2015	18	16
Treasury stock, at cost - 247,566,270 shares as of June 30, 2016 and 247,566,270 shares as of December 31, 2015	(1,717)	(1,717)
Additional paid-in capital	16,923	16,860
Accumulated deficit	(8,934)	(8,927)
Accumulated other comprehensive income (loss), net of tax	(70)	88
Total stockholders' equity	6,220	6,320
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY	$17,838	$18,133

See notes to the unaudited condensed consolidated financial statements.

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BOSTON SCIENTIFIC CORPORATION AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS (UNAUDITED)

	Six Months Ended June 30,	
in millions	2016	2015
Cash provided by (used for) operating activities	$537	$(137)
Investing activities:		
Purchases of property, plant and equipment	(138)	(92)
Proceeds from disposal of property, plant and equipment	29	—
Purchases of privately-held securities	(36)	(140)
Purchases of notes receivable	(5)	—
Payments for acquisitions of businesses, net of cash acquired	—	(63)
Payments for investments and acquisitions of certain technologies	—	(2)
Cash provided by (used for) investing activities	(150)	(297)
Financing activities:		
Payments on long-term borrowings	(250)	(1,000)
Proceeds from long-term borrowings, net of debt issuance costs	—	1,831
Payment of contingent consideration	(35)	(87)
Proceeds from borrowings on credit facilities	40	395
Payments on borrowings from credit facilities	(40)	(395)
Cash used to net share settle employee equity awards	(56)	(62)
Proceeds from issuances of shares of common stock	73	70
Cash provided by (used for) financing activities	(268)	752
Effect of foreign exchange rates on cash	—	(2)
Net increase (decrease) in cash and cash equivalents	119	316
Cash and cash equivalents at beginning of period	319	587
Cash and cash equivalents at end of period	$438	$903
Supplemental Information		
Stock-based compensation expense	$58	$53
Fair value of contingent consideration recorded in purchase accounting	—	31

See notes to the unaudited condensed consolidated financial statements.

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NOTES TO THE CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (UNAUDITED)

NOTE A – BASIS OF PRESENTATION

The accompanying unaudited condensed consolidated financial statements of Boston Scientific Corporation have been prepared in accordance with accounting principles generally accepted in the United States (U.S. GAAP) and with the instructions to Form 10-Q and Article 10 of Regulation S-X. Accordingly, they do not include all of the information and footnotes required by U.S. GAAP for complete financial statements. In the opinion of management, all adjustments (consisting only of normal recurring adjustments) considered necessary for fair presentation have been included. Operating results for the three and six month periods ended June 30, 2016 are not necessarily indicative of the results that may be expected for the year ending December 31, 2016. For further information, refer to the consolidated financial statements and footnotes thereto included in Item 8 of our most recent Annual Report on Form 10-K.

Subsequent Events

We evaluate events occurring after the date of our most recent accompanying unaudited condensed consolidated balance sheets for potential recognition or disclosure in our financial statements. We did not identify any material subsequent events requiring adjustment to our accompanying unaudited condensed consolidated financial statements (recognized subsequent events) for the three and six month periods ended June 30, 2016. Those items requiring disclosure (unrecognized subsequent events) in the financial statements have been disclosed accordingly. Refer to Note B - Acquisitions and Strategic Investments, Note H - Income Taxes and Note I - Commitments and Contingencies for more information.

Pension Termination Charges

Following our 2006 acquisition of Guidant Corporation, we sponsored the Guidant Retirement Plan, a frozen noncontributory defined benefit plan covering a select group of current and former employees. The plan was partially frozen as of September 25, 1995 and completely frozen as of May 31, 2007. The plan was subsequently terminated effective December 1, 2014. During 2015, we finalized the termination process and settled the plan’s obligations, and as a result, we recorded pension termination charges of $8 million during the first half of 2015 and an additional $36 million during the third quarter of 2015 for a total of $44 million of pension termination charges in the year ended December 31, 2015. We do not expect to record any additional pension termination charges in 2016 related to the termination of the Guidant Retirement Plan.

NOTE B – ACQUISITIONS AND STRATEGIC INVESTMENTS

2016 Acquisitions

We did not close any material acquisitions during the first half of 2016.

On July 27, 2016, we acquired Cosman Medical, Inc. (Cosman), a privately held manufacturer of radiofrequency ablation systems, expanding our Neuromodulation portfolio and offering physicians treating patients with chronic pain a wider choice of non-opioid therapeutic options. We plan to begin the process of integrating Cosman into our Neuromodulation business during the second half of 2016.

2015 Acquisitions

Xlumena, Inc.

On April 2, 2015, we acquired Xlumena, Inc. (Xlumena), a medical device company that developed minimally invasive devices for Endoscopic Ultrasound (EUS) guided transluminal drainage of targeted areas within the gastrointestinal tract. The purchase agreement called for an up-front payment of $63 million, an additional payment of $13 million upon FDA clearance of the HOT AXIOS™ product, and further sales-based milestones based on sales achieved through 2018. We are in the process of integrating Xlumena into our Endoscopy business, and expect the integration to be substantially complete by the end of 2016.

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Purchase Price Allocation

We accounted for the acquisition of Xlumena as a business combination and, in accordance with Financial Accounting Standards Board (FASB) Accounting Standards Codification® (ASC) Topic 805, Business Combinations, we have recorded the assets acquired and liabilities assumed at their respective fair values as of the acquisition date. The components of the aggregate purchase price are as follows (in millions):

Cash, net of cash acquired	$63
Fair value of contingent consideration	31
	$94

The following summarizes the aggregate purchase price allocation for the 2015 acquisition as of June 30, 2015 (in millions):

Goodwill	$30
Amortizable intangible assets	68
Inventory	1
Other net assets	2
Deferred income taxes	(7)
	$94

We allocated a portion of the purchase price to specific intangible asset categories as follows:

	Amount Assigned (in millions)	Weighted Average Amortization Period (in years)	Range of Risk-Adjusted Discount Rates used in Purchase Price Allocation
Amortizable intangible assets:			
Technology-related	$ 68	11	15 %
	$ 68		

Contingent Consideration

Certain of our acquisitions involve contingent consideration arrangements. Payment of additional consideration is generally contingent on the acquired company reaching certain performance milestones, including attaining specified revenue levels, achieving product development targets and/or obtaining regulatory approvals. In accordance with U.S. GAAP, we recognize a liability equal to the fair value of the contingent payments we expect to make as of the acquisition date. We re-measure this liability each reporting period and record changes in the fair value through a separate line item within our condensed consolidated statements of operations.

We recorded net expenses related to the changes in fair value of our contingent consideration liabilities of $33 million during the second quarter of 2016, $37 million during the first half of 2016, $19 million during the second quarter of 2015 and $46 million during the first half of 2015. We paid contingent consideration of $14 million during the second quarter of 2016, $77 million during the first half of 2016, $11 million during the second quarter of 2015 and $110 million during the first half of 2015.

Changes in the fair value of our contingent consideration liabilities were as follows (in millions):

Balance as of December 31, 2015	$246
Other amounts recorded related to prior acquisitions	1
Fair value adjustments	37
Contingent payments related to prior period acquisitions	(77)
Balance as of June 30, 2016	$207

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As of June 30, 2016, the maximum amount of future contingent consideration (undiscounted) that we could be required to pay was approximately $1.572 billion.

Contingent consideration liabilities are remeasured to fair value each reporting period using projected revenues, discount rates, probabilities of payment and projected payment dates. The recurring Level 3 fair value measurements of our contingent consideration liabilities include the following significant unobservable inputs:

Contingent Consideration Liabilities	Fair Value as of June 30, 2016	Valuation Technique	Unobservable Input	Range
R&D, regulatory and commercialization-based Milestones	$17 million	Discounted Cash Flow	Discount Rate	2.0% - 2.9%
			Probability of Payment	0% - 64%
			Projected Year of Payment	2018 - 2021
Revenue-based Payments	$41 million	Discounted Cash Flow	Discount Rate	14% - 15%
			Projected Year of Payment	2017 - 2020
	$149 million	Monte Carlo	Revenue Volatility	20%
			Risk Free Rate	LIBOR Term Structure & Cost of Debt Structure
			Projected Year of Payment	2016 - 2022

Increases or decreases in the fair value of our contingent consideration liabilities can result from changes in discount periods and rates, as well as changes in the timing and amount of revenue estimates or in the timing or likelihood of achieving R&D, regulatory commercialization-based and revenue-based milestones. Projected contingent payment amounts related to some of our R&D, regulatory and commercialization-based and revenue-based milestones are discounted back to the current period using a discounted cash flow (DCF) model. Other revenue-based payments are valued using a Monte Carlo valuation model, which simulates future revenues during the earn-out period using management's best estimates. Projected revenues are based on our most recent internal operational budgets and long-range strategic plans. Increases in projected revenues and probabilities of payment may result in higher fair value measurements. Increases in discount rates and the time to payment may result in lower fair value measurements. Increases or decreases in any of those inputs together, or in isolation, may result in a significantly lower or higher fair value measurement.

Strategic Investments

We did not close any material strategic investments during the first half of 2016.

On April 30, 2015, we acquired a 27 percent ownership interest in Preventice, Inc. (Preventice), which includes 18.5 percent of Preventice's common stock. Preventice is a privately-held company headquartered in Minneapolis, MN, and a leading developer of mobile health solutions and services. Preventice offers a full portfolio of wearable cardiac monitors, including Holter monitors, cardiac event monitors and mobile cardiac telemetry. In addition to the equity agreement, we entered into a commercial agreement with Preventice, under which we have become Preventice’s exclusive, worldwide sales and marketing representative. We believe this partnership strengthens our portfolio of cardiac monitoring and broader disease management capabilities.

On April 13, 2015, we acquired 25 percent of the common stock of Frankenman Medical Equipment Company (Frankenman). Frankenman is a privately-held company headquartered in Suzhou, China, and is a local market leader in surgical staplers. Additionally, we entered into co-promotional and co-selling agreements with Frankenman to

jointly commercialize selected products in China. We believe this alliance will enable us to reach more clinicians and treat more patients in China by providing access to training on less invasive endoscopic technologies with clinical and economic benefits.

We account for certain of our strategic investments as equity method investments, in accordance with Financial Accounting Standards Board (FASB) Accounting Standards Codification (ASC) Topic 323, Investments - Equity Method and Joint Ventures. The book value of investments that we accounted for under the equity method of accounting was $221 million as of June 30, 2016 and $173 million as of December 31, 2015. The aggregate value of our cost method investments was $25 million as of June 30, 2016 and $45 million as of December 31, 2015. In addition we had notes receivable from certain companies that we account for under the cost method of $32 million as of June 30, 2016 and $30 million as of December 31, 2015.

As of June 30, 2016, the book value of our equity method investments exceeded our share of the book value of the investees' underlying net assets by approximately $140 million, which represents amortizable intangible assets and in-process research and

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development, corresponding deferred tax liabilities, and goodwill. During the three and six months ended June 30, 2016 and June 30, 2015, the net losses from our equity method adjustments, presented within the Other, net caption of our condensed consolidated statement of operations were immaterial.

NOTE C – GOODWILL AND OTHER INTANGIBLE ASSETS

The gross carrying amount of goodwill and other intangible assets and the related accumulated amortization for intangible assets subject to amortization and accumulated write-offs of goodwill as of June 30, 2016 and December 31, 2015 are as follows:

	As of June 30, 2016		December 31, 2015	
(in millions)	Gross Carrying Amount	Accumulated Amortization/ Write-offs	Gross Carrying Amount	Accumulated Amortization/ Write-offs
Amortizable intangible assets				
Technology-related	$8,948	$ (4,257)	$8,948	$ (4,054)
Patents	522	(366)	520	(358)
Other intangible assets	1,531	(666)	1,529	(610)
	$11,001	$ (5,289)	$10,997	$ (5,022)
Unamortizable intangible assets				
Goodwill	$16,375	$ (9,900)	$16,373	$ (9,900)
In-process research and development (IPR&D)	98	—	99	—
Technology-related	120	—	120	—
	$16,593	$ (9,900)	$16,592	$ (9,900)

Our technology-related intangible assets that are not subject to amortization represent technical processes, intellectual property and/or institutional understanding acquired through business combinations that are fundamental to the on-going operations of our business and have no limit to their useful life. Our technology-related intangible assets that are not subject to amortization are comprised primarily of certain acquired balloon and other technology, which is foundational to our continuing operations within the Cardiovascular market and other markets within interventional medicine. We assess our indefinite-lived intangible assets at least annually for impairment and reassess their classification as indefinite-lived assets. We assess qualitative factors to determine whether the existence of events and circumstances indicate that it is more likely than not that our indefinite-lived intangible assets are impaired. If we conclude that it is more likely than not that the asset is impaired, we then determine the fair value of the intangible asset and perform the quantitative impairment test by comparing the fair value with the carrying value in accordance with ASC Topic 350, Intangibles - Goodwill and Other.

The following represents our goodwill balance by global reportable segment:

(in millions)	Cardiovascular	Rhythm Management	MedSurg	Total
Balance as of December 31, 2015	$ 3,451	$ 292	$ 2,730	$6,473
Purchase price adjustments	—	—	2	2
Balance as of June 30, 2016	$ 3,451	$ 292	$ 2,732	$6,475

Goodwill Impairment Testing

In the second quarter of 2016, we performed our annual goodwill impairment test for all of our reporting units and concluded the fair value of each reporting unit exceeded its carrying value. Based on the criteria prescribed in FASB ASC Topic 350 - Intangibles - Goodwill and Other, we assess goodwill for impairment at the reporting unit level,

which is defined as an operating segment or one level below an operating segment, referred to as a component. In 2016 and 2015, we identified six operating segments including Interventional Cardiology, Peripheral Interventions, Rhythm Management, Endoscopy, Urology and Pelvic Health, and Neuromodulation. For purposes of identifying our reporting units, we then assessed whether any components of these segments constitute a business for which discrete financial information is available and where segment management regularly reviews the operating results of that component. We identified Rhythm Management as having two components: Cardiac Rhythm Management and Electrophysiology.

For our 2016 and 2015 annual impairment assessment, we identified seven reporting units, which align to our seven core businesses: Interventional Cardiology, Peripheral Interventions, Cardiac Rhythm Management, Electrophysiology, Endoscopy, Urology and

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Pelvic Health, and Neuromodulation. For our 2016 annual impairment assessment the Cardiac Rhythm Management and Electrophysiology reporting units, components of the Rhythm Management operating segment, were aggregated due to a reorganization commencing in 2015 which resulted in integrated leadership, shared resources and consolidation of certain sites in 2016. Because our global Electrophysiology reporting unit was identified as being at higher risk of potential goodwill impairment during our 2015 annual test, it was tested for impairment on a stand-alone basis in the second quarter of 2016, immediately prior to aggregating it with our global Cardiac Rhythm Management reporting unit. The fair value exceeded the carrying value by approximately 36 percent. In comparison, the global Electrophysiology reporting unit had excess fair value of approximately 28 percent as of our 2015 annual test.

As of the date of our 2016 annual goodwill impairment test, the aggregated global Electrophysiology and Cardiac Rhythm Management reporting unit (Rhythm Management) had excess fair value over carrying value of approximately 70 percent and held $292 million of allocated goodwill. As such, it was not deemed at higher risk of future impairment. Changes in our reporting units or in the structure of our business as a result of future reorganizations, acquisitions or divestitures of assets or businesses could result in future impairments of goodwill within our reporting units. Refer to Critical Accounting Policies and Estimates within our Management's Discussion and Analysis of Financial Condition and Results of Operations contained in Item 2 of this Quarterly Report on Form 10-Q for a discussion of key assumptions used in our testing.

On a quarterly basis, we monitor the key drivers of fair value to detect events or other changes that would warrant an interim impairment test of our goodwill. The key variables that drive the cash flows of our reporting units and amortizable intangibles are estimated revenue growth rates and levels of profitability. Terminal value growth rate assumptions, as well as the Weighted Average Cost of Capital (WACC) rate applied are additional key variables for reporting unit cash flows. These assumptions are subject to uncertainty, including our ability to grow revenue and improve profitability levels. The estimates used for our future cash flows and discount rates represent management's best estimates, which we believe to be reasonable, but future declines in business performance may impair the recoverability of our goodwill.

Future events that could have a negative impact on the levels of excess fair value over carrying value of our reporting units include, but are not limited to:

- decreases in estimated market sizes or market growth rates due to greater-than-expected declines in procedural volumes, pricing pressures, reductions in reimbursement levels, product actions, and/or competitive technology developments;

- declines in our market share and penetration assumptions due to increased competition, an inability to develop or launch new and next-generation products and technology features in line with our commercialization strategies, and market and/or regulatory conditions that may cause significant launch delays or product recalls;

- decreases in our forecasted profitability due to an inability to successfully implement and achieve timely and sustainable cost improvement measures consistent with our expectations;

- negative developments in intellectual property litigation that may impact our ability to market certain products or increase our costs to sell certain products;

- the level of success of ongoing and future research and development efforts, including those related to recent acquisitions, and increases in the research and development costs necessary to obtain regulatory approvals and launch new products;

- the level of success in managing the growth of acquired companies, achieving sustained profitability consistent with our expectations, establishing government and third-party payer reimbursement, supplying the market, and increases in the costs and time necessary to integrate acquired businesses into our operations successfully;

- changes in our reporting units or in the structure of our business as a result of future reorganizations, acquisitions or divestitures of assets or businesses; and

- increases in our market-participant risk-adjusted WACC, and increases in our market-participant tax rate, and/or changes in tax laws or macroeconomic conditions.

Negative changes in one or more of these factors, among others, could result in impairment charges.

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The following is a rollforward of accumulated goodwill write-offs by global reportable segment:

(in millions)	Cardiovascular	Rhythm Management	MedSurg	Total
Accumulated write-offs as of December 31, 2015	$ (1,479)	$ (6,960)	$(1,461)	$(9,900)
Goodwill written off	—	—	—	—
Accumulated write-offs as of June 30, 2016	$ (1,479)	$ (6,960)	$(1,461)	$(9,900)

Intangible Asset Impairment Testing

On a quarterly basis, we monitor for events or other potential indicators of impairment that would warrant an interim impairment test of our intangible assets. We did not record any intangible asset impairment charges during the six months ended June 30, 2016.

2015 Charges

During the second quarter of 2015, in conjunction with our annual strategic planning process and annual goodwill impairment test, we performed an interim impairment test on certain of our IPR&D projects and core technology assets. Based on our impairment assessment, we recorded an impairment charge of $9 million in the second quarter of 2015.

The nonrecurring Level 3 fair value measurements of our intangible asset impairment analysis included the following significant unobservable inputs:

Intangible Asset	Valuation Date	Fair Value	Valuation Technique	Unobservable Input	Rate
In-Process R&D	June 30, 2015	$6 million	Income Approach - Excess Earnings Method	Discount Rate	16.5 - 20%

NOTE D – FAIR VALUE MEASUREMENTS

Derivative Instruments and Hedging Activities

We address market risk from changes in foreign currency exchange rates and interest rates through a risk management program that includes the use of derivative financial instruments, and we operate the program pursuant to documented corporate risk management policies. Our derivative instruments do not subject our earnings or cash flows to material risk, as gains and losses on these derivatives generally offset losses and gains on the item being hedged. We do not enter into derivative transactions for speculative purposes, and we do not have any non-derivative instruments that are designated as hedging instruments pursuant to FASB ASC Topic 815, Derivatives and Hedging (Topic 815).

Currency Hedging

We are exposed to currency risk consisting primarily of foreign currency denominated monetary assets and liabilities, forecasted foreign currency denominated intercompany and third-party transactions and net investments in certain subsidiaries. We manage our exposure to changes in foreign currency exchange rates on a consolidated basis to take advantage of offsetting transactions. We use derivative instruments and non-derivative transactions to reduce the risk that our earnings and cash flows associated with these foreign currency denominated balances and transactions will be adversely affected by foreign currency exchange rate changes.

Currently or Previously Designated Foreign Currency Hedges

All of our designated currency hedge contracts outstanding as of June 30, 2016 and December 31, 2015 were cash flow hedges under Topic 815 intended to protect the U.S. dollar value of our forecasted foreign currency denominated transactions. We record the effective portion of any change in the fair value of foreign currency cash flow hedges in other comprehensive income (OCI) until the related third-party transaction occurs. Once the related third-party transaction occurs, we reclassify the effective portion of any related gain or loss on the foreign currency cash flow hedge to earnings. In the event the hedged forecasted transaction does not occur, or it becomes no longer probable that it will occur, we reclassify the amount of any gain or loss on the related cash flow hedge to earnings at that time. We had currency derivative instruments designated as cash flow hedges outstanding in the contract amount of $2.476 billion as of June 30, 2016 and $1.458 billion as of December 31, 2015.

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We recognized net gains of $32 million in earnings on our cash flow hedges during the second quarter of 2016 and $80 million for the first half of 2016, as compared to net gains of $53 million during the second quarter of 2015 and $102 million for the first half of 2015. All currency cash flow hedges outstanding as of June 30, 2016 mature within 60 months. As of June 30, 2016, $7 million of net loss, net of tax, was recorded in accumulated other comprehensive income (AOCI) to recognize the effective portion of the fair value of any currency derivative instruments that are, or previously were, designated as foreign currency cash flow hedges, as compared to net gains, net of tax, of $145 million as of December 31, 2015. As of June 30, 2016, $49 million of net gains, net of tax, may be reclassified to earnings within the next twelve months.

The success of our hedging program depends, in part, on forecasts of transaction activity in various currencies (primarily British pound sterling, Euro and Japanese yen). We may experience unanticipated currency exchange gains or losses to the extent that there are differences between forecasted and actual activity during periods of currency volatility. In addition, changes in foreign currency exchange rates related to any unhedged transactions may impact our earnings and cash flows.

Non-designated Foreign Currency Contracts

We use currency forward contracts as a part of our strategy to manage exposure related to foreign currency denominated monetary assets and liabilities. These currency forward contracts are not designated as cash flow, fair value or net investment hedges under Topic 815; are marked-to-market with changes in fair value recorded to earnings; and are entered into for periods consistent with currency transaction exposures, generally less than one year. We had currency derivative instruments not designated as hedges under Topic 815 outstanding in the contract amount of $2.389 billion as of June 30, 2016 and $2.090 billion as of December 31, 2015.

Interest Rate Hedging

Our interest rate risk relates primarily to U.S. dollar borrowings, partially offset by U.S. dollar cash investments. We have historically used interest rate derivative instruments to manage our earnings and cash flow exposure to changes in interest rates by converting floating-rate debt into fixed-rate debt or fixed-rate debt into floating-rate debt. We had no interest rate derivative instruments outstanding as of June 30, 2016.

We designate these derivative instruments either as fair value or cash flow hedges under Topic 815. We record changes in the value of fair value hedges in interest expense, which is generally offset by changes in the fair value of the hedged debt obligation. Interest payments made or received related to our interest rate derivative instruments are included in interest expense. We record the effective portion of any change in the fair value of derivative instruments designated as cash flow hedges as unrealized gains or losses in OCI, net of tax, until the hedged cash flow occurs, at which point the effective portion of any gain or loss is reclassified to earnings. We record the ineffective portion of our cash flow hedges in interest expense. In the event the hedged cash flow does not occur, or it becomes no longer probable that it will occur, we reclassify the amount of any gain or loss on the related cash flow hedge to interest expense at that time.

In the fourth quarter of 2013, we entered into interest rate derivative contracts having a notional amount of $450 million to convert fixed-rate debt into floating-rate debt, which we designated as fair value hedges. During the first quarter of 2015, we terminated these hedges, and we received total proceeds of approximately $35 million, which included approximately $7 million of net accrued interest receivable. We assessed at inception, and re-assessed on an ongoing basis, whether the interest rate derivative contracts were highly effective in offsetting changes in the fair value of the hedged fixed-rate debt. We recognized no gains or losses in interest expense, related to fair value hedges, during the second quarter of 2015. During the first half of 2015, we recognized, in interest expense, an $8 million loss

on our hedged debt and an $8 million gain on the related interest rate derivative contract.

We are amortizing the gains and losses on previously terminated interest rate derivative instruments, including fixed-to-floating interest rate contracts designated as fair value hedges, and forward starting interest rate derivative contracts and treasury locks designated as cash flow hedges upon termination into earnings as a component of interest expense over the remaining term of the hedged debt, in accordance with Topic 815. The carrying amount of certain of our senior notes included unamortized gains of $57 million as of June 30, 2016 and $63 million as of December 31, 2015. We had immaterial unamortized losses as of June 30, 2016 and December 31, 2015 related to the fixed-to-floating interest rate contracts. In addition, we had pre-tax net gains within AOCI related to terminated forward starting interest rate derivative contracts and treasury locks of $9 million as of June 30, 2016 and $10 million as of December 31, 2015. The net gains that we recognized as a reduction of interest expense in earnings related to previously terminated interest rate derivatives were approximately $3 million during the second quarter of 2016 and $6 million during the first half of 2016, as compared to $5 million during the second quarter of 2015 and $7 million during the first half of 2015. As of June 30, 2016, $13 million of pre-tax net gains may be reclassified to earnings within the next twelve months as a reduction to interest expense from amortization of our terminated interest rate derivative contracts.

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Counterparty Credit Risk

We do not have significant concentrations of credit risk arising from our derivative financial instruments, whether from an individual counterparty or a related group of counterparties. We manage the concentration of counterparty credit risk on our derivative instruments by limiting acceptable counterparties to a diversified group of major financial institutions with investment grade credit ratings, limiting the amount of credit exposure to each counterparty, and actively monitoring their credit ratings and outstanding fair values on an ongoing basis. Furthermore, none of our derivative transactions are subject to collateral or other security arrangements, and none contain provisions that are dependent on our credit ratings from any credit rating agency.

We also employ master netting arrangements that reduce our counterparty payment settlement risk on any given maturity date to the net amount of any receipts or payments due between us and the counterparty financial institution. Thus, the maximum loss due to counterparty credit risk is limited to the unrealized gains in such contracts net of any unrealized losses should any of these counterparties fail to perform as contracted. Although these protections do not eliminate concentrations of credit risk, as a result of the above considerations, we do not consider the risk of counterparty default to be significant.

Fair Value of Derivative Instruments

The following presents the effect of our derivative instruments designated as cash flow hedges under Topic 815 on our accompanying unaudited condensed consolidated statements of operations during the second quarter and first half of 2016 and 2015 (in millions):

	Amount of Pre-tax Gain (Loss) Recognized in OCI (Effective Portion)	Amount of Pre-tax Gain (Loss) Reclassified from AOCI into Earnings (Effective Portion)	Location in Statement of Operations
Three Months Ended June 30, 2016			
Currency hedge contracts	$ (99)	$ 32	Cost of products sold
	$ (99)	$ 32	
Three Months Ended June 30, 2015			
Currency hedge contracts	$ (25)	$ 53	Cost of products sold
Interest rate derivative contracts	$ 10	$ 1	Interest Expense
	$ (15)	$ 54	
Six Months Ended June 30, 2016			
Currency hedge contracts	$ (158)	$ 80	Cost of products sold
	$ (158)	$ 80	
Six Months Ended June 30, 2015			
Currency hedge contracts	$ 68	$ 102	Cost of products sold
Interest rate derivative contracts	$ 11	$ 2	Interest Expense
	$ 79	$ 104	

The amount of gain (loss) recognized in earnings related to the ineffective portion of hedging relationships was immaterial for all periods presented.

Net gains and losses on currency hedge contracts not designated as hedging instruments were offset by net losses and gains from foreign currency transaction exposures, as shown in the following table:

in millions	Location in Statement of Operations	Three Months Ended June 30,		Six Months Ended June 30,	
		2016	2015	2016	2015
Gain (loss) on currency hedge contracts	Other, net	$(28)	$(9)	$(67)	$14
Gain (loss) on foreign currency transaction exposures	Other, net	29	4	63	(28)
Net foreign currency gain (loss)	Other, net	$1	$(5)	$(4)	$(14)

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Topic 815 requires all derivative instruments to be recognized at their fair values as either assets or liabilities on the balance sheet. We determine the fair value of our derivative instruments using the framework prescribed by FASB ASC Topic 820, Fair Value Measurements and Disclosures (Topic 820), by considering the estimated amount we would receive or pay to transfer these instruments at the reporting date and by taking into account current interest rates, foreign currency exchange rates, the creditworthiness of the counterparty for the assets and our creditworthiness for liabilities. In certain instances, we may utilize financial models to measure fair value. In doing so, we use inputs that include quoted prices for similar assets or liabilities in active markets; quoted prices for identical or similar assets or liabilities in markets that are not active; other observable inputs for the asset or liability; and inputs derived principally from, or corroborated by, observable market data by correlation or other means. As of June 30, 2016, we have classified all of our derivative assets and liabilities within Level 2 of the fair value hierarchy prescribed by Topic 820, as discussed below, because these observable inputs are available for substantially the full term of our derivative instruments.

The following are the balances of our derivative assets and liabilities as of June 30, 2016 and December 31, 2015:

		As of	
		June 30,	December 31,
(in millions)	Location in Balance Sheet (1)	2016	2015
Derivative Assets:			
Currently or Previously Designated Hedging Instruments			
Currency hedge contracts	Other current assets	$88	$ 138
Currency hedge contracts	Other long-term assets	13	66
		101	204
Non-Designated Hedging Instruments			
Currency hedge contracts	Other current assets	43	33
Total Derivative Assets		$144	$ 237
Derivative Liabilities:			
Currently or Previously Designated Hedging Instruments			
Currency hedge contracts	Other current liabilities	$21	$ 1
Currency hedge contracts	Other long-term liabilities	99	—
		120	1
Non-Designated Hedging Instruments			
Currency hedge contracts	Other current liabilities	79	22
Total Derivative Liabilities		$199	$ 23

(1) We classify derivative assets and liabilities as current when the remaining term of the derivative contract is one year or less.

Other Fair Value Measurements

Recurring Fair Value Measurements

On a recurring basis, we measure certain financial assets and financial liabilities at fair value based upon quoted market prices, where available. Where quoted market prices or other observable inputs are not available, we apply valuation techniques to estimate fair value. Topic 820 establishes a three-level valuation hierarchy for disclosure of fair value measurements. The categorization of financial assets and financial liabilities within the valuation hierarchy is based upon the lowest level of input that is significant to the measurement of fair value. The three levels of the hierarchy are defined as follows:

- Level 1 – Inputs to the valuation methodology are quoted market prices for identical assets or liabilities.

- Level 2 – Inputs to the valuation methodology are other observable inputs, including quoted market prices for similar assets or liabilities and market-corroborated inputs.

- Level 3 – Inputs to the valuation methodology are unobservable inputs based on management's best estimate of inputs market participants would use in pricing the asset or liability at the measurement date, including assumptions about risk.

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Assets and liabilities measured at fair value on a recurring basis consist of the following as of June 30, 2016 and December 31, 2015:

	June 30, 2016				As of December 31, 2015			
(in millions)	Level 1	Level 2	Level 3	Total	Level 1	Level 2	Level 3	Total
Assets								
Money market and government funds	$77	$—	$—	$77	$118	$—	$—	$118
Currency hedge contracts	—	144	—	144	—	237	—	237
	$77	$144	$—	$221	$118	$237	$—	$355
Liabilities								
Currency hedge contracts	$—	$199	$—	$199	$—	$23	$—	$23
Accrued contingent consideration	—	—	207	207	—	—	246	246
	$—	$199	$207	$406	$—	$23	$246	$269

Our investments in money market and government funds are classified within Level 1 of the fair value hierarchy because they are valued using quoted market prices. These investments are classified as cash and cash equivalents within our accompanying unaudited condensed consolidated balance sheets, in accordance with U.S. GAAP and our accounting policies. In addition to $77 million invested in money market and government funds as of June 30, 2016, we had $60 million in short-term time deposits and $301 million in interest bearing and non-interest bearing bank accounts. In addition to $118 million invested in money market and government funds as of December 31, 2015, we had $31 million in short-term deposits and $170 million in interest bearing and non-interest bearing bank accounts.

Our recurring fair value measurements using significant unobservable inputs (Level 3) relate solely to our contingent consideration liabilities. Refer to Note B - Acquisitions and Strategic Investments for a discussion of the changes in the fair value of our contingent consideration liabilities.

Non-Recurring Fair Value Measurements

We hold certain assets and liabilities that are measured at fair value on a non-recurring basis in periods subsequent to initial recognition. The fair value of a cost method investment is not estimated if there are no identified events or changes in circumstances that may have a significant adverse effect on the fair value of the investment. Refer to Note B – Acquisitions and Strategic Investments for a discussion of our strategic investments.

The fair value of our outstanding debt obligations was $5.878 billion as of June 30, 2016 and $5.887 billion as of December 31, 2015, which was determined by using primarily quoted market prices for our publicly registered senior notes, classified as Level 1 within the fair value hierarchy. Refer to Note E – Borrowings and Credit Arrangements for a discussion of our debt obligations.

NOTE E – BORROWINGS AND CREDIT ARRANGEMENTS

We had total debt of $5.427 billion as of June 30, 2016 and $5.677 billion as of December 31, 2015. The debt maturity schedule for the significant components of our debt obligations as of June 30, 2016 is as follows:

(in millions)	2016	2017	2018	2019	2020	Thereafter	Total
Senior Notes	$—	$250	$600	$—	$1,450	$ 2,350	$4,650
Term Loans	—	—	225	150	375	—	750
	$—	$250	$825	$150	$1,825	$ 2,350	$5,400

Note: The table above does not include unamortized discounts associated with our senior notes, or amounts related to interest rate contracts used to hedge the fair value of certain of our senior notes.

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Revolving Credit Facility

On April 10, 2015, we entered into a new $2.000 billion revolving credit facility (the 2015 Facility) with a global syndicate of commercial banks and terminated our previous $2.000 billion revolving credit facility. The 2015 Facility matures on April 10, 2020. Eurodollar and multicurrency loans under the 2015 Facility bear interest at LIBOR plus an interest margin of between 0.900 percent and 1.500 percent, based on our corporate credit ratings and consolidated leverage ratio (1.300 percent as of June 30, 2016). In addition, we are required to pay a facility fee based on our credit ratings, consolidated leverage ratio and the total amount of revolving credit commitment, regardless of usage, under the credit agreement (0.200 percent per year as of June 30, 2016). The 2015 Facility contains covenants which, among other things, require that we maintain a minimum interest coverage ratio of 3.0 times consolidated EBITDA and a maximum leverage ratio of 4.5 times consolidated EBITDA for the first four fiscal quarter-ends following the closing of the acquisition of the American Medical Systems male urology portfolio (AMS Portfolio Acquisition) on August 3, 2015, and decreasing to 4.25 times, 4.0 times, and 3.75 times consolidated EBITDA for the next three fiscal quarter-ends after such four fiscal quarter-ends, respectively, and then to 3.50 times for each fiscal quarter-end thereafter. There were no amounts borrowed under our current and prior revolving credit facilities as of June 30, 2016 or December 31, 2015.

	Covenant Requirement as of June 30, 2016	Actual as of June 30, 2016
Maximum leverage ratio (1)	4.5 times	2.6 times
Minimum interest coverage ratio (2)	3.0 times	8.8 times

(1) Ratio of total debt to consolidated EBITDA, as defined by the credit agreement, for the preceding four consecutive fiscal quarters.

(2) Ratio of consolidated EBITDA, as defined by the credit agreement, to interest expense for the preceding four consecutive fiscal quarters.

The credit agreement for the 2015 Facility provides for an exclusion from the calculation of consolidated EBITDA, as defined by the credit agreement, through the credit agreement maturity, of any non-cash charges and up to $620 million in restructuring charges and restructuring-related expenses related to our current or future restructuring plans. As of June 30, 2016, we had $523 million of the restructuring charge exclusion remaining. In addition, any cash litigation payments (net of any cash litigation receipts), as defined by the agreement, are excluded from the calculation of consolidated EBITDA and any new debt issued to fund any tax deficiency payments is excluded from consolidated total debt, as defined in the agreement, provided that the sum of any excluded net cash litigation payments and any new debt issued to fund any tax deficiency payments not exceed $2.000 billion in the aggregate. As of June 30, 2016, we had $1.543 billion of the combined legal and debt exclusion remaining.

As of and through June 30, 2016, we were in compliance with the required covenants.

Term Loans

As of June 30, 2016, we had an aggregate of $750 million outstanding under our unsecured term loan facilities and $1.000 billion outstanding as of December 31, 2015. These facilities include an unsecured term loan facility entered into in August 2013 (2013 Term Loan) which had $150 million outstanding as of June 30, 2016 and $250 million outstanding as of December 31, 2015, along with an unsecured term loan credit facility entered into in April 2015 (2015 Term Loan) which had $600 million outstanding as of June 30, 2016 and $750 million outstanding as of December 31, 2015.

Borrowings under the 2013 Term Loan bear interest at LIBOR plus an interest margin between 1.00 percent and 1.75 percent (currently 1.50 percent) based on our corporate credit ratings and consolidated leverage ratio. We repaid $150 million of our 2013 Term Loan facility in the fourth quarter of 2015 and repaid an additional $100 million during the second quarter of 2016. As a result and in accordance with the credit agreement, the outstanding balance of $150 million is the remaining principal amount due at the final maturity date in August 2018. The 2013 Term Loan borrowings are repayable at any time without premium or penalty. Our term loan facility requires that we comply with certain covenants, including financial covenants with respect to maximum leverage and minimum interest coverage, consistent with the 2015 Term Loan facility. The maximum leverage ratio requirement is 4.5 times, and our actual leverage ratio as of June 30, 2016 is 2.6 times. The minimum interest coverage ratio requirement is 3.0 times, and our actual interest coverage ratio as of June 30, 2016 is 8.8 times.

On April 10, 2015, we entered into a new $750 million unsecured term loan credit facility (2015 Term Loan) which matures on August 3, 2020. The 2015 Term Loan was funded on August 3, 2015 and was used to partially fund the AMS Portfolio Acquisition, including the payment of fees and expenses. Term loan borrowings under this facility bear interest at LIBOR plus an interest margin of between 1.00 percent and 1.75 percent (currently 1.50 percent), based on our corporate credit ratings and consolidated

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leverage ratio. We repaid $150 million of our 2015 Term Loan during the second quarter of 2016. The remaining 2015 Term Loan requires quarterly principal payments of $38 million commencing in the third quarter of 2018, and the remaining principal amount is due at the final maturity date of August 3, 2020. The 2015 Term Loan agreement requires that we comply with certain covenants, including financial covenants with respect to maximum leverage and minimum interest coverage, consistent with our revolving credit facility. The maximum leverage ratio requirement is 4.5 times, and our actual leverage ratio as of June 30, 2016 is 2.6 times. The minimum interest coverage ratio requirement is 3.0 times, and our actual interest coverage ratio as of June 30, 2016 is 8.8 times.

Senior Notes

We had senior notes outstanding of $4.650 billion as of June 30, 2016 and December 31, 2015. In May 2015, we completed the offering of $1.850 billion in aggregate principal amount of senior notes consisting of $600 million in aggregate principal amount of 2.850% notes due 2020, $500 million in aggregate principal amount of 3.375% notes due 2022 and $750 million in aggregate principal amount of 3.850% notes due 2025. The net proceeds from the offering of the notes, after deducting underwriting discounts and estimated offering expenses, were approximately $1.831 billion. We used a portion of the net proceeds from the senior notes offering to redeem $400 million aggregate principal amount of our 5.500% notes due November 2015 and $600 million aggregate principal amount of our 6.400% notes due June 2016. The remaining senior notes offering proceeds, together with the 2015 Term Loan, were used to fund the AMS Portfolio Acquisition. We recorded a charge of $45 million in interest expense, during the second quarter of 2015, for premiums, accelerated amortization of debt issuance costs, and investor discount costs net of interest rate hedge gains related to the early debt extinguishment.

Our senior notes were issued in public offerings, are redeemable prior to maturity and are not subject to any sinking fund requirements. Our senior notes are unsecured, unsubordinated obligations and rank on parity with each other. These notes are effectively junior to borrowings under our credit and security facility, to the extent if borrowed by our subsidiaries and to liabilities of our subsidiaries (see Other Arrangements below).

Other Arrangements

We maintain a $300 million credit and security facility secured by our U.S. trade receivables maturing on June 9, 2017. The credit and security facility requires that we maintain a maximum leverage covenant consistent with our revolving credit facility. The maximum leverage ratio requirement is 4.5 times, and our actual leverage ratio as of June 30, 2016 is 2.6 times. We had no borrowings outstanding under this facility as of June 30, 2016 and December 31, 2015.

We have accounts receivable factoring programs in certain European countries that we account for as sales under FASB ASC Topic 860, Transfers and Servicing. These agreements provide for the sale of accounts receivable to third parties, without recourse, of up to approximately $400 million as of June 30, 2016. We have no retained interests in the transferred receivables, other than collection and administrative responsibilities and, once sold, the accounts receivable are no longer available to satisfy creditors in the event of bankruptcy. We de-recognized $169 million of receivables as of June 30, 2016 at an average interest rate of 1.8 percent, and $151 million as of December 31, 2015 at an average interest rate of 2.4 percent.

In addition, we have uncommitted credit facilities with a commercial Japanese bank that provide for borrowings, promissory notes discounting and receivables factoring of up to 21.000 billion Japanese yen (approximately $204 million as of June 30, 2016). We de-recognized $170 million of notes receivable and factored receivables as of June 30, 2016 at an average interest rate of 1.6 percent and $132 million of notes receivable as of December 31, 2015 at an average interest rate of 1.6 percent. De-recognized accounts and notes receivable are excluded from trade accounts receivable, net in the accompanying unaudited condensed consolidated balance sheets.

As of June 30, 2016 we had outstanding letters of credit of $43 million, as compared to $44 million as of December 31, 2015, which consisted primarily of bank guarantees and collateral for workers' compensation insurance arrangements. As of June 30, 2016 and December 31, 2015, none of the beneficiaries had drawn upon the letters of credit or guarantees; accordingly, we did not recognize a related liability for our outstanding letters of credit in our consolidated balance sheets as of June 30, 2016 or December 31, 2015. We believe we will generate sufficient cash from operations to fund these arrangements and intend to fund these arrangements without drawing on the letters of credit.

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NOTE F – RESTRUCTURING-RELATED ACTIVITIES

On an ongoing basis, we monitor the dynamics of the economy, the healthcare industry, and the markets in which we compete. We continue to assess opportunities for improved operational effectiveness and efficiency, and better alignment of expenses with revenues, while preserving our ability to make the investments in research and development projects, capital and our people that we believe are essential to our long-term success. As a result of these assessments, we have undertaken various restructuring initiatives in order to enhance our growth potential and position us for long-term success. These initiatives are described below.

2016 Restructuring Plan

On June 6, 2016, our Board of Directors approved, and we committed to, a restructuring initiative (the 2016 Restructuring Plan). The 2016 Restructuring Plan is intended to develop global commercialization, technology and manufacturing capabilities in key growth markets, build on our Plant Network Optimization (PNO) strategy which is intended to simplify our manufacturing plant structure by transferring certain production lines among facilities, and expand operational efficiencies in support of our operating income margin goals. Key activities under the 2016 Restructuring Plan include strengthening global infrastructure through evolving global real estate and workplaces, developing global commercial and technical competencies, enhancing manufacturing and distribution expertise in certain regions, and continuing implementation of our ongoing PNO strategy. These activities initiated in the second quarter of 2016 and are expected to be substantially completed by the end of 2018.

The implementation of the 2016 Restructuring Plan is expected to result in total pre-tax charges of approximately $175 million to $225 million, and approximately $160 million to $210 million of these charges are estimated to result in cash outlays, of which we have made payments of $5 million through June 30, 2016. We have recorded related costs of $19 million since the inception of the plan through June 30, 2016, and recorded a portion of these expenses as restructuring charges and the remaining portion through other lines within our consolidated statements of operations.

The following table provides a summary of our estimates of costs associated with the 2016 Restructuring Plan through the end of 2018 by major type of cost:

Type of cost	Total estimated amount expected to be incurred
Restructuring charges:	
Termination benefits	$65 million to $80 million
Other (1)	$15 million to $25 million
Restructuring-related expenses:	
Other (2)	$95 million to $120 million
	$175 million to $225 million

(1) Consists primarily of consulting fees and costs associated with contract cancellations.
(2) Comprised of other costs directly related to the 2016 Restructuring Plan, including program management, accelerated depreciation, and costs to transfer product lines among facilities.

2014 Restructuring Plan

On October 22, 2013, our Board of Directors approved, and we committed to, a restructuring initiative (the 2014 Restructuring Plan). The 2014 Restructuring Plan is intended to build on the progress we have made to address financial pressures in a changing global marketplace, further strengthen our operational effectiveness and efficiency and support new growth investments. Key activities under the plan include continued implementation of our ongoing PNO strategy, continued focus on driving operational efficiencies and ongoing business and commercial model changes. The PNO strategy is intended to simplify our manufacturing plant structure by transferring certain

production lines among facilities. Other activities involve rationalizing organizational reporting structures to streamline various functions, eliminate bureaucracy, increase productivity and better align resources to business strategies and marketplace dynamics. These activities were initiated in the fourth quarter of 2013 and were substantially completed by the end of 2015, except for certain ongoing actions associated with our PNO strategy, which we expect to be substantially completed by the end of 2016.

The implementation of the 2014 Restructuring Plan is expected to result in total pre-tax charges of approximately $255 million to $270 million, and approximately $240 million to $255 million of these charges are estimated to result in cash outlays, of which we have made payments of $224 million through June 30, 2016. We have recorded related costs of $249 million since the inception of the plan, and recorded a portion of these expenses as restructuring charges and the remaining portion through other lines within our consolidated statements of operations.

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The following table provides a summary of our estimates of costs associated with the 2014 Restructuring Plan through the end of 2016 by major type of cost:

Type of cost	Total estimated amount expected to be incurred
Restructuring charges:	
Termination benefits	$95 million to $100 million
Other (1)	$30 million to $35 million
Restructuring-related expenses:	
Other (2)	$130 million to $135 million
	$255 million to $270 million

(1) Consists primarily of consulting fees and costs associated with contract cancellations.
(2) Comprised of other costs directly related to the 2014 Restructuring Plan, including program management, accelerated depreciation, and costs to transfer product lines among facilities.

We recorded net restructuring charges pursuant to our restructuring plans of $14 million in the second quarter of 2016, $3 million in the second quarter of 2015, $17 million in the first half of 2016 and $9 million in the first half of 2015. In addition, we recorded expenses within other lines of our accompanying unaudited condensed consolidated statements of operations related to our restructuring initiatives of $12 million in the second quarter of 2016, $12 million in the second quarter of 2015, $22 million in the first half of 2016 and $28 million in the first half of 2015.

The following presents these costs (credits) by major type and line item within our accompanying unaudited condensed consolidated statements of operations, as well as by program:

Three Months Ended June 30, 2016

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
Restructuring charges	$ 14	$ —	$ —	$ —	$ 14
Restructuring-related expenses:					
Cost of products sold	—	—	7	—	7
Selling, general and administrative expenses	—	3	—	2	5
	—	3	7	2	12
	$ 14	$ 3	$ 7	$ 2	$ 26

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
2016 Restructuring Plan	$ 18	$ —	$ 1	$ —	$ 19
2014 Restructuring Plan	(4)	3	6	2	7
	$ 14	$ 3	$ 7	$ 2	$ 26

Three Months Ended June 30, 2015

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
Restructuring charges	$ 3	$ —	$ —	$ —	$ 3
Restructuring-related expenses:					
Cost of products sold	—	—	8	—	8
Selling, general and administrative expenses	—	1	—	3	4
	—	1	8	3	12
	$ 3	$ 1	$ 8	$ 3	$ 15

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All charges incurred in the second quarter of 2015 were related to the 2014 Restructuring Plan.

Six Months Ended June 30, 2016

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
Restructuring charges	$ 15	$ —	$ —	$ 2	$17
Restructuring-related expenses:					
Cost of products sold	—	—	12	—	12
Selling, general and administrative expenses	—	4	—	6	10
	—	4	12	6	22
	$ 15	$ 4	$ 12	$ 8	$39

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
2016 Restructuring Plan	$ 18	$ —	$ 1	$ —	$19
2014 Restructuring Plan	(3)	4	11	8	20
	$ 15	$ 4	$ 12	$ 8	$39

Six Months Ended June 30, 2015

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
Restructuring charges	$ 8	$ —	$ —	$ 1	$9
Restructuring-related expenses:					
Cost of products sold	—	—	16	—	16
Selling, general and administrative expenses	—	2	—	10	12
	—	2	16	10	28
	$ 8	$ 2	$ 16	$ 11	$37

(in millions)	Termination Benefits	Accelerated Depreciation	Transfer Costs	Other	Total
2014 Restructuring Plan	$ 11	$ 2	$ 16	$ 11	$40
Substantially completed restructuring programs	(3)	—	—	—	(3)
	$ 8	$ 2	$ 16	$ 11	$37

Termination benefits represent amounts incurred pursuant to our ongoing benefit arrangements and amounts for "one-time" involuntary termination benefits, and have been recorded in accordance with FASB ASC Topic 712, Compensation – Non-retirement Postemployment Benefits and FASB ASC Topic 420, Exit or Disposal Cost Obligations (Topic 420). We expect to record additional termination benefits related to our 2016 Restructuring Plan throughout the rest of 2016 when we identify with more specificity the job classifications, functions and locations of headcount reductions. We do not expect to record any additional termination benefits related to our 2014 Restructuring Plan. Other restructuring costs, which represent primarily consulting fees and costs related to contract cancellations, are being recorded as incurred in accordance with Topic 420. Accelerated depreciation is being recorded over the adjusted remaining useful life of the related assets, and program management and production line transfer costs are being recorded as incurred.

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As of June 30, 2016, we incurred cumulative restructuring charges related to our 2016 Restructuring Plan and our 2014 Restructuring Plan of $142 million and restructuring-related costs of $126 million since we committed to the plans. The following presents these costs by major type:

(in millions)	2016 Restructuring Plan	2014 Restructuring Plan	Total
Termination benefits	$ 18	$ 93	$111
Other	—	31	31
Total restructuring charges	18	124	142
Accelerated depreciation	—	12	12
Transfer costs	1	66	67
Other	—	47	47
Restructuring-related expenses	1	125	126
	$ 19	$ 249	$268

We made cash payments of $17 million in the second quarter of 2016 and $40 million in the first half of 2016 associated with our restructuring initiatives, and as of June 30, 2016, we had made total cash payments of $229 million related to our 2016 Restructuring Plan and 2014 Restructuring Plan since committing to the plans. These payments were made using cash generated from operations, and are comprised of the following:

(in millions)	2016 Restructuring Plan	2014 Restructuring Plan	Total
Three Months Ended June 30, 2016			
Termination benefits	$ 4	$ 3	$7
Transfer costs	1	6	7
Other	—	3	3
	$ 5	$ 12	$17
Six Months Ended June 30, 2016			
Termination benefits	$ 4	$ 17	$21
Transfer costs	1	11	12
Other	—	7	7
	$ 5	$ 35	$40
Program to Date			
Termination benefits	$ 4	$ 86	$90
Transfer costs	1	66	67
Other	—	72	72
	$ 5	$ 224	$229

Our restructuring liability is primarily comprised of accruals for termination benefits. The following is a rollforward of the termination benefit liability associated with our 2016 Restructuring Plan and our 2014 Restructuring Plan, which is reported as a component of accrued expenses included in our accompanying unaudited condensed balance sheets:

(in millions)	2016 Restructuring Plan	2014 Restructuring Plan	Total
Accrued as of December 31, 2015	$ —	$ 29	$29

Charges (credits)	18	(3)	15
Cash payments	(4)	(17)	(21)
Accrued as of June 30, 2016	$ 14	$ 9	$ 23

In addition to our accrual for termination benefits, we had a $4 million liability as of June 30, 2016 and a $3 million liability as of December 31, 2015 for other restructuring-related items.

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NOTE G – SUPPLEMENTAL BALANCE SHEET INFORMATION

Components of selected captions in our accompanying unaudited condensed consolidated balance sheets are as follows:

Trade accounts receivable, net

(in millions)	As of June 30, 2016	December 31, 2015
Accounts receivable	$1,512	$ 1,394
Less: allowance for doubtful accounts	(80)	(75)
Less: allowance for sales returns	(45)	(44)
	$1,387	$ 1,275

The following is a rollforward of our allowance for doubtful accounts for the second quarter and first half of 2016 and 2015:

	Three Months Ended June 30,		Six Months Ended June 30,	
(in millions)	2016	2015	2016	2015
Beginning balance	$80	$72	$75	$76
Charges to expenses	3	6	7	8
Utilization of allowances	(3)	(1)	(2)	(7)
Ending balance	$80	$77	$80	$77

Inventories

(in millions)	As of June 30, 2016	December 31, 2015
Finished goods	$654	$ 706
Work-in-process	108	102
Raw materials	219	208
	$981	$ 1,016

Property, plant and equipment, net

(in millions)	As of June 30, 2016	December 31, 2015
Land	$83	$ 86
Buildings and improvements	966	981
Equipment, furniture and fixtures	2,904	2,793
Capital in progress	183	202
	4,136	4,062
Less: accumulated depreciation	2,649	2,572
	$1,487	$ 1,490

Depreciation expense was $62 million for the second quarter of 2016, $65 million for the second quarter of 2015, $126 million for the first half of 2016, and $130 million for the first half of 2015.

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Accrued expenses

(in millions)	As of June 30, 2016	December 31, 2015
Legal reserves	$1,075	$ 773
Payroll and related liabilities	500	504
Accrued contingent consideration	102	119
Other	559	574
	$2,236	$ 1,970

Other long-term liabilities

(in millions)	As of June 30, 2016	December 31, 2015
Accrued income taxes	$1,291	$ 1,253
Legal reserves	1,300	1,163
Accrued contingent consideration	105	127
Other long-term liabilities	543	431
	$3,239	$ 2,974

Accrued warranties

We offer warranties on certain of our product offerings. The majority of our warranty liability relates to implantable devices offered by our Cardiac Rhythm Management (CRM) business, which include defibrillator and pacemaker systems. Our CRM products come with a standard limited warranty covering the replacement of these devices. We offer a full warranty for a portion of the period post-implant, and a partial warranty for a period of time thereafter. We estimate the costs that we may incur under our warranty programs based on the number of units sold, historical and anticipated rates of warranty claims and cost per claim, and record a liability equal to these estimated costs as cost of products sold at the time the product sale occurs. We reassess the adequacy of our recorded warranty liabilities on a quarterly basis and adjust these amounts as necessary. The current portion of our warranty accrual is included in other accrued expenses in the table above and the non-current portion of our warranty accrual is included in other long-term liabilities in the table above. Changes in our product warranty accrual during the first half of 2016 and 2015 consisted of the following (in millions):

	Six Months Ended June 30, 2016	2015
Beginning Balance	$23	$25
Provision	10	10
Settlements/reversals	(13)	(9)
Ending Balance	$20	$26

NOTE H – INCOME TAXES

Our effective tax rates from continuing operations for the three months ended June 30, 2016 and June 30, 2015, were 47.8% and 2.9%, respectively. For the first half of 2016 and 2015 our effective tax rates from continuing operations were 97.0% and (86.9)%, respectively. The change in our reported tax rate for the second quarter and first half of 2016, as compared to the same periods in 2015, relates primarily to the impact of certain receipts and charges that are

taxed at different rates than our effective tax rate, including acquisition-related items, contingent consideration, litigation-related and restructuring-related items, as well as the impact of certain discrete tax items.

As of June 30, 2016, we had $1.058 billion of gross unrecognized tax benefits, of which a net $902 million, if recognized, would affect our effective tax rate. As of December 31, 2015, we had $1.056 billion of gross unrecognized tax benefits, of which a net $900 million, if recognized, would affect our effective tax rate.

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We have received Notices of Deficiency from the Internal Revenue Service (IRS) reflecting proposed audit adjustments for Guidant Corporation for its 2001 through 2006 tax years and for Boston Scientific Corporation for its 2006 and 2007 tax years. The total incremental tax liability asserted by the IRS for the applicable periods is $1.162 billion plus interest. The primary issue in dispute for all years is the transfer pricing associated with the technology license agreements between domestic and foreign subsidiaries of Guidant. In addition, the IRS has proposed adjustments in connection with the financial terms of our Transaction Agreement with Abbott Laboratories pertaining to the sale of Guidant's vascular intervention business to Abbott in April 2006. During 2014, we received a Revenue Agent Report from the IRS reflecting significant proposed audit adjustments to our 2008, 2009, and 2010 tax years based upon the same transfer pricing methodologies that the IRS applied to our 2001 through 2007 tax years.

We do not agree with the transfer pricing methodologies applied by the IRS or its resulting assessments. We have filed petitions with the U.S. Tax Court (Tax Court) contesting the Notices of Deficiency for the 2001 through 2007 tax years in challenge and submitted a letter to the IRS Office of Appeals (IRS Appeals) protesting the Revenue Agent Report for the 2008 through 2010 tax years and requesting an administrative appeal hearing. The issues in dispute were scheduled to be heard in Tax Court in late July 2016. On July 19, 2016, we entered into a Stipulation of Settled Issues with the IRS intended to resolve all of the aforementioned transfer pricing issues, as well as the issues related to our transaction with Abbott. The Stipulation of Settled Issues is contingent upon IRS Appeals applying the same basis of settlement to all transfer pricing issues for the Company’s 2008, 2009, and 2010 tax years, and if applicable, review by the U.S. Congress Joint Committee on Taxation.

In the event that the conditions in the Stipulation of Settled Items are satisfied, we expect to make net tax payments to the IRS of approximately $275 million, plus interest through the date of payment. No payments will be required until the dispute, including its resolution with IRS Appeals, is definitively resolved and our tax liability, including interest, for each individual year has been recalculated by the IRS. Based on experiences of other companies, we expect to make payment within the next 12 to 24 months. We believe our income tax reserves associated with these matters are adequate as of June 30, 2016 and do not expect any additional charges related to the resolution of this controversy.

We recognize interest and penalties related to income taxes as a component of income tax expense. We had $535 million accrued for gross interest and penalties as of June 30, 2016 and $500 million as of December 31, 2015. We recognized net tax expense related to interest and penalties of $13 million during the second quarter of 2016, $10 million during the second quarter of 2015, $23 million during the first half of 2016 and $21 million during the first half of 2015.

It is reasonably possible that within the next 12 months we will resolve multiple issues including transfer pricing and transactional-related issues with foreign, federal and state taxing authorities, in which case we could record a reduction in our balance of unrecognized tax benefits of up to approximately $767 million.

In November 2015, the FASB issued ASC Update No. 2015-17, Income Taxes (Topic 740): Balance Sheet Classification of Deferred Taxes. This update simplifies the presentation of deferred income taxes by requiring all deferred tax assets and liabilities, along with any related valuation allowance, to be classified as noncurrent on the balance sheet. The new guidance is effective for all public Companies for annual periods beginning after December 15, 2016, and interim periods within those annual periods, with early adoption permitted. We elected to early adopt this standard prospectively at the beginning of 2016.

NOTE I – COMMITMENTS AND CONTINGENCIES

The medical device market in which we primarily participate is largely technology driven. As a result, intellectual property rights, particularly patents and trade secrets, play a significant role in product development and differentiation. Over the years, there has been litigation initiated against us by others, including our competitors,

claiming that our current or former product offerings infringe patents owned or licensed by them. Intellectual property litigation is inherently complex and unpredictable. In addition, competing parties frequently file multiple suits to leverage patent portfolios across product lines, technologies and geographies and to balance risk and exposure between the parties. In some cases, several competitors are parties in the same proceeding, or in a series of related proceedings, or litigate multiple features of a single class of devices. These forces frequently drive settlement not only for individual cases, but also for a series of pending and potentially related and unrelated cases. Although monetary and injunctive relief is typically sought, remedies and restitution are generally not determined until the conclusion of the trial court proceedings and can be modified on appeal. Accordingly, the outcomes of individual cases are difficult to time, predict or quantify and are often dependent upon the outcomes of other cases in other geographies.

During recent years, we successfully negotiated closure of several long-standing legal matters and have received favorable rulings in several other matters; however, there continues to be outstanding intellectual property litigation. Adverse outcomes in one or more of these matters could have a material adverse effect on our ability to sell certain products and on our operating margins, financial position, results of operations and/or liquidity.

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In the normal course of business, product liability, securities and commercial claims are asserted against us. Similar claims may be asserted against us in the future related to events not known to management at the present time. We maintain an insurance policy providing limited coverage against securities claims, and we are substantially self-insured with respect to product liability claims and fully self-insured with respect to intellectual property infringement claims. The absence of significant third-party insurance coverage increases our potential exposure to unanticipated claims or adverse decisions. Product liability claims, securities and commercial litigation, and other legal proceedings in the future, regardless of their outcome, could have a material adverse effect on our financial position, results of operations and/or liquidity.

In addition, like other companies in the medical device industry, we are subject to extensive regulation by national, state and local government agencies in the United States and other countries in which we operate. From time to time we are the subject of qui tam actions and governmental investigations often involving regulatory, marketing and other business practices. These qui tam actions and governmental investigations could result in the commencement of civil and criminal proceedings, substantial fines, penalties and administrative remedies and have a material adverse effect on our financial position, results of operations and/or liquidity.

In accordance with ASC Topic 450, Contingencies, we accrue anticipated costs of settlement, damages, losses for product liability claims and, under certain conditions, costs of defense, based on historical experience or to the extent specific losses are probable and estimable. Otherwise, we expense these costs as incurred. If the estimate of a probable loss is a range and no amount within the range is more likely, we accrue the minimum amount of the range.

Our accrual for legal matters that are probable and estimable was $2.375 billion as of June 30, 2016 and $1.936 billion as of December 31, 2015, and includes certain estimated costs of settlement, damages and defense. We recorded $628 million of litigation-related charges during the first half of 2016 and $192 million of litigation-related charges during the first half of 2015. We continue to assess certain litigation and claims to determine the amounts, if any, that management believes will be paid as a result of such claims and litigation and, therefore, additional losses may be accrued and paid in the future, which could materially adversely impact our operating results, cash flows and/or our ability to comply with our debt covenants.

In management's opinion, we are not currently involved in any legal proceedings other than those disclosed in our most recent Annual Report on Form 10-K and our Quarterly Report on Form 10-Q for the quarter ended March 31, 2016 and those specifically identified below, which, individually or in the aggregate, could have a material adverse effect on our financial condition, operations and/or cash flows. Unless included in our legal accrual or otherwise indicated below, a range of loss associated with any individual material legal proceeding cannot be estimated.

Patent Litigation

On April 19, 2016, a subsidiary of Boston Scientific filed suit against Edwards Lifesciences Corporation in the United States District Court for the District of Delaware for patent infringement. We allege that Edwards' SAPIEN 3 valve infringes a patent related to adaptive sealing technology. On June 9, 2016, Edwards filed a counterclaim alleging that our Lotus™ transcatheter heart valve system infringes three patents owned by Edwards.

Product Liability Litigation

As of August 1, 2016, over 39,000 product liability cases or claims related to transvaginal surgical mesh products designed to treat stress urinary incontinence and pelvic organ prolapse have been asserted against us. The pending cases are in various federal and state courts in the United States and include eight putative class actions. There were also fewer than 20 cases in Canada, inclusive of four putative class actions, and fewer than 20 claims in the United Kingdom. Generally, the plaintiffs allege personal injury associated with use of our transvaginal surgical mesh

products. The plaintiffs assert design and manufacturing claims, failure to warn, breach of warranty, fraud, violations of state consumer protection laws and loss of consortium claims. Over 3,100 of the cases have been specially assigned to one judge in state court in Massachusetts. On February 7, 2012, the Judicial Panel on Multi-District Litigation (MDL) established MDL-2326 in the U.S. District Court for the Southern District of West Virginia and transferred the federal court transvaginal surgical mesh cases to MDL-2326 for coordinated pretrial proceedings. During the fourth quarter of 2013, we received written discovery requests from certain state attorneys general offices regarding our transvaginal surgical mesh products. We have responded to those requests. As of August 1, 2016, we have entered into master settlement agreements with certain plaintiffs' counsel to resolve an aggregate of approximately 11,000 cases and claims. These master settlement agreements provide that the settlement and distribution of settlement funds to participating claimants are conditional upon, among other things, achieving minimum required claimant participation thresholds. Of the 11,000 cases and claims, 6,000 have met the conditions of the settlement and are final. All settlement agreements were entered into solely by way of compromise and without any admission or concession by us of any liability or wrongdoing. In addition, we continue to engage in discussions with various

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plaintiffs' counsel regarding potential resolution of pending cases and claims and, as of August 1, 2016, have made substantial progress in discussions with plaintiffs' counsel representing approximately 8,000 additional cases and claims.

We have established a product liability accrual for known and estimated future cases and claims asserted against us as well as with respect to the actions that have resulted in verdicts against us and the costs of defense thereof associated with our transvaginal surgical mesh products. While we believe that our accrual associated with this matter is adequate, changes to this accrual may be required in the future as additional information becomes available. While we continue to engage in discussions with plaintiffs' counsel regarding potential resolution of pending cases and claims and intend to vigorously contest the cases and claims asserted against us; that do not settle, the final resolution of the cases and claims is uncertain and could have a material impact on our results of operations, financial condition and/or liquidity. Initial trials involving our transvaginal surgical mesh products have resulted in both favorable and unfavorable judgments for us. We do not believe that the judgment in any one trial is representative of potential outcomes of all cases or claims related to our transvaginal surgical mesh products.

Governmental Investigations and Qui Tam Matters

On May 5, 2014, we were served with a subpoena from the U.S. Department of Health and Human Services, Office of the Inspector General. The subpoena seeks information relating to the launch of the Cognis and Teligen line of devices in 2008, the performance of those devices from 2007 to 2009, and the operation of the Physician Guided Learning Program. On May 6, 2016, a qui tam lawsuit in this matter was unsealed in the U.S. District Court for the District of Minnesota. At the same time, we learned that U.S. Government and the State of California had earlier declined to intervene in that lawsuit on April 15, 2016. The complaint was served on us on July 21, 2016

Other Proceedings

On September 28, 2011, we served a complaint against Mirowski Family Ventures LLC in the U.S. District Court for the Southern District of Indiana for a declaratory judgment that we have paid all royalties owed and did not breach any contractual or fiduciary obligations arising out of a license agreement. Mirowski answered and filed counterclaims requesting damages. On May 13, 2013, Mirowski Family Ventures served us with a complaint alleging breach of contract in Montgomery County Circuit Court, Maryland, and they amended this complaint on August 1, 2013. On July 29, 2013, the Indiana case was dismissed. On September 10, 2013, we removed the case to the United States District Court for the District of Maryland. On June 5, 2014, the District Court granted Mirowski's motion to remand the case to the Montgomery County Circuit Court. On September 24, 2014, following a jury verdict against us, the Montgomery County Circuit Court entered a judgment that we breached our license agreement with Mirowski and awarded damages of $308 million. On October 28, 2014, the Montgomery County Circuit Court denied our post-trial motions seeking to overturn the judgment. On November 19, 2014, we filed an appeal with the Maryland Court of Special Appeals. On January 29, 2016, the Maryland Court of Special Appeals affirmed the decision of the Montgomery County Circuit Court. On February 2, 2016, we filed a motion for reconsideration, which was denied. On July 12, 2016, the Maryland Court of Appeals denied our petition for certiorari. We plan to seek United States Supreme Court review. On July 26, 2016, we paid $366 million in satisfaction of the judgment and interest, subject to a right of rescission should the judgment be reversed.
Refer to Note H - Income Taxes for information regarding our tax litigation.

Matters Concluded Since December 31, 2015

On April 24, 2014, Dr. Qingsheng Zhu and Dr. Julio Spinelli, acting jointly on behalf of the stockholder representative committee of Action Medical, Inc. (Action Medical), filed a lawsuit against us and our subsidiary, Cardiac Pacemakers, Inc. (CPI), in the U.S. District Court for the District of Delaware. The stockholder representatives alleged

that we and CPI breached a contractual duty to pursue development and commercialization of certain patented heart pacing methods and devices and to return certain patents. On March 15, 2016, the Court granted summary judgment in our favor as to all of plaintiffs' claims for damages. The parties subsequently reached a resolution on the remaining claim and counterclaim concerning specific performance, and the case was dismissed on June 29, 2016.

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NOTE J – WEIGHTED AVERAGE SHARES OUTSTANDING

	Three Months Ended June 30,		Six Months Ended June 30,	
(in millions)	2016	2015	2016	2015
Weighted average shares outstanding - basic	1,357.4	1,341.3	1,353.9	1,337.5
Net effect of common stock equivalents	—	*20.5	—	*22.2
Weighted average shares outstanding - assuming dilution	1,357.4	1,361.8	1,353.9	1,359.7

*We generated net losses in the second quarter and first half of 2016. Our weighted-average shares outstanding for earnings per share calculations exclude common stock equivalents of 17.7 million for the second quarter of 2016 and 18.6 million for the first half of 2016 due to our net loss positions.

Weighted average shares outstanding, assuming dilution, excludes the impact of three million stock options for the second quarter of 2015 and three million for the first half of 2015, due to the exercise prices of these stock options being greater than the average fair market value of our common stock during the period. This impact was immaterial for the second quarter and first half of 2016.

We issued approximately four million shares of our common stock in the second quarter of 2016, two million shares of our common stock in the second quarter of 2015, 12 million shares of our common stock in the first half of 2016 and15 million shares of our common stock in the first half of 2015, following the exercise of underlying stock options or vesting of deferred stock units, or purchases under our employee stock purchase plan. We did not repurchase any shares of our common stock during the first half of 2016 or 2015.

NOTE K – SEGMENT REPORTING

In 2016 and 2015, we identified six operating segments including Interventional Cardiology, Peripheral Interventions, Rhythm Management, Endoscopy, Urology and Pelvic Health, and Neuromodulation. Our reportable segments represent an aggregate of operating segments based on the criteria prescribed in FASB ASC Topic 280 - Segment Reporting. We have three reportable segments comprised of Cardiovascular, Rhythm Management, and MedSurg. Each of our reportable segments generates revenues from the sale of medical devices. We measure and evaluate our reportable segments based on segment net sales and operating income, excluding the impact of changes in foreign currency. Sales generated from reportable segments, as well as operating results of reportable segments, are based on internally-derived standard currency exchange rates, which may differ from year to year, and do not include intersegment profits. As needed, we restate segment information for the prior period based on our internally-derived standard currency exchange rates used for the current period in order to remove the impact of foreign currency exchange fluctuation. We exclude from segment operating income certain corporate-related expenses and certain charges or credits that our chief operating decision maker considers to be non-recurring and/or non-operational, such as amounts related to intangible asset impairment charges, acquisition- and divestiture-, litigation-, restructuring- and restructuring-related net charges and credits; pension termination charges; and amortization expense. Although we exclude these amounts from segment operating income, they are included in reported consolidated operating income (loss) and are included in the reconciliation below.

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A reconciliation of the totals reported for the reportable segments to the applicable line items in our accompanying unaudited condensed consolidated statements of operations is as follows:

(in millions)	Three Months Ended June 30, 2016	2015	Six Months Ended June 30, 2016	2015
Net sales				
Interventional Cardiology	$635	$567	$1,248	$1,108
Peripheral Interventions	278	245	542	477
Cardiovascular	913	812	1,790	1,585
Cardiac Rhythm Management	512	490	983	973
Electrophysiology	64	60	128	121
Rhythm Management	576	550	1,111	1,094
Endoscopy	390	352	755	680
Urology and Pelvic Health	270	142	513	272
Neuromodulation	139	125	264	241
MedSurg	799	619	1,532	1,193
Net sales allocated to reportable segments	2,288	1,981	4,433	3,872
Impact of foreign currency fluctuations	(162)	(138)	(343)	(261)
	$2,126	$1,843	$4,090	$3,611
Income (loss) before income taxes				
Cardiovascular	$294	$247	$593	$483
Rhythm Management	97	78	187	155
MedSurg	262	188	502	355
Operating income allocated to reportable segments	653	513	1,282	993
Corporate expenses and currency exchange	(155)	(105)	(290)	(188)
Intangible asset impairment charges; acquisition- and divestiture-, restructuring- and restructuring-related net charges; litigation-related net charges and credits; and pension termination charges	(697)	(73)	(762)	(333)
Amortization expense	(135)	(116)	(271)	(229)
Operating income (loss)	(334)	219	(41)	243
Other expense, net	(63)	(114)	(128)	(189)
Income (loss) before income taxes	$(397)	$105	$(169)	$54

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NOTE L – CHANGES IN OTHER COMPREHENSIVE INCOME

The following table provides the reclassifications out of other comprehensive income for the three and six months ended June 30, 2016 and June 30, 2015. Amounts in the chart below are presented net of tax.

Three Months Ended June 30, 2016

(in millions)	Foreign currency translation adjustments	Unrealized gains/losses on derivative financial instruments	Defined benefit pension items / Other	Total
Balance as of March 31, 2016	$ (38)	$ 83	$ (10)	$35
Other comprehensive income (loss) before reclassifications	(21)	(64)	(1)	(86)
Amounts reclassified from accumulated other comprehensive income	—	(20)	1	(19)
Net current-period other comprehensive income	(21)	(84)	—	(105)
Balance as of June 30, 2016	$ (59)	$ (1)	$ (10)	$(70)

Three Months Ended June 30, 2015

(in millions)	Foreign currency translation adjustments	Unrealized gains/losses on derivative financial instruments	Defined benefit pension items / Other	Total
Balance as of March 31, 2015	$ (73)	$ 247	$ (32)	$142
Other comprehensive income (loss) before reclassifications	5	(8)	(2)	(5)
Amounts reclassified from accumulated other comprehensive income	—	(35)	2	(33)
Net current-period other comprehensive income	5	(43)	—	(38)
Balance as of June 30, 2015	$ (68)	$ 204	$ (32)	$104

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Six Months Ended June 30, 2016

(in millions)	Foreign currency translation adjustments	Unrealized gains/losses on derivative financial instruments	Defined benefit pension items / Other	Total
Balance as of December 31, 2015	$ (54)	$ 152	$ (10)	$88
Other comprehensive income (loss) before reclassifications	(5)	(102)	(3)	(110)
Amounts reclassified from accumulated other comprehensive income	—	(51)	3	(48)
Net current-period other comprehensive income	(5)	(153)	—	(158)
Balance as of June 30, 2016	$ (59)	$ (1)	$ (10)	$(70)

Six Months Ended June 30, 2015

(in millions)	Foreign currency translation adjustments	Unrealized gains/losses on derivative financial instruments	Defined benefit pension items / Other	Total
Balance as of December 31, 2014	$ (38)	$ 219	$ (37)	$144
Other comprehensive income (loss) before reclassifications	(30)	51	(4)	17
Amounts reclassified from accumulated other comprehensive income	—	(66)	9	(57)
Net current-period other comprehensive income	(30)	(15)	5	(40)
Balance as of June 30, 2015	$ (68)	$ 204	$ (32)	$104

The income tax impact of the amounts in other comprehensive income for unrealized gains and losses on derivative financial instruments before reclassifications was a benefit of $36 million in the second quarter of 2016, a benefit of $5 million in the second quarter of 2015, a benefit of $57 million in the first half of 2016 and an expense of $28 million in the first half of 2015. The gains and losses on derivative financial instruments reclassified were reduced by income tax impacts of $12 million in the second quarter of 2016, $19 million in the second quarter of 2015, $29 million in the first half of 2016 and $37 million in the first half of 2015. Refer to Note D – Fair Value Measurements in this Quarterly Report on Form 10-Q for further detail on the reclassifications related to derivatives.

The income tax impact of the amounts in other comprehensive income for defined benefit and pension items before reclassification was an immaterial benefit in both the second quarter of 2016 and the second quarter of 2015. For both the first half of 2016 and the first half of 2015, the income tax impact of the amounts in other comprehensive income for defined benefit and pension items before reclassification was immaterial.
The gains and losses on defined benefit and pension related items reclassified from accumulated other comprehensive income were reduced by immaterial income tax impacts in the second quarter and first half of 2016. The gains and losses on defined benefit and pension related items reclassified from accumulated other comprehensive income were reduced by immaterial income tax impacts in the second quarter of 2015 and by $5 million in the first half of 2015.

NOTE M – NEW ACCOUNTING PRONOUNCEMENTS

From time to time, new accounting pronouncements are issued by the FASB or other standard setting bodies. Recently issued standards typically do not require adoption until a future effective date. Prior to their effective date, we evaluate the pronouncements to determine the potential effects of adoption on our consolidated financial statements.

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Standards Implemented

ASC Update No. 2015-05

In April 2015, the FASB issued ASC Update No. 2015-05, Intangibles- Goodwill and Other - Internal -Use Software (Subtopic 350-40): Customer's Accounting for Fees Paid in a Cloud Computing Arrangement. Update No. 2015-05 provides accounting guidance on how customers should treat cloud computing arrangements. If a cloud computing arrangement includes a software license, then the customer should account for the software license element of the arrangement consistent with the acquisition of other software licenses. If a cloud computing arrangement does not include a software license, the customer should account for the arrangement as a service contract. Update No. 2015-05 is effective for annual reporting periods beginning after December 15, 2015 and interim periods within those reporting periods. We elected to adopt the amendments prospectively to all arrangements entered into or materially modified after the effective date. The adoption of Update No. 2015-05 did not have a material impact on our financial position or results of operations.

ASC Update No. 2015-12

In July 2015, the FASB issued ASC Update No. 2015-12, Plan Accounting: Defined Benefit Pension Plans (Topic 960), Defined Contribution Pension Plans (Topic 962), and Health and Welfare Benefit Plans (Topic 965). Update No. 2015-12 has three parts. Part I designates contract value as the only required measure for fully benefit-responsive investment contracts. Part II simplifies the investment disclosure requirements under Topics 820, 960, 962, and 965 for employee benefits plans and Part III provides an alternative measurement date for fiscal periods that do not coincide with a month-end date. Update No. 2015-12 is effective for fiscal years beginning after December 15, 2015. The adoption of Update No. 2015-12 did not have a material impact on our financial position or results of operations.

ASC Update No. 2015-16

In September 2015, the FASB issued ASC Update No. 2015-16, Business Combinations (Topic 805): Simplifying the Accounting for Measurement - Period Adjustments. Update No. 2015-16 eliminates the requirement to restate prior period financial statements for measurement period adjustments following a business combination. Update No. 2015-16 requires that the cumulative impact of a measurement period adjustment (including the impact on prior periods) be recognized in the reporting period in which the adjustment is identified. The prior period impact of the adjustment should be either presented separately on the face of the income statement or disclosed in the notes. Update No. 2015-16 is effective for fiscal years beginning after December 15, 2015. The adoption of Update No. 2015-16 did not impact on our financial position or results of operations.

ASC Update No. 2015-17

In November 2015, the FASB issued ASC Update No. 2015-17, Income Taxes (Topic 740): Balance Sheet Classification of Deferred Taxes. Update No. 2015-17 requires that deferred tax liabilities and assets be classified as noncurrent in a classified statement of financial position. It is effective for fiscal years beginning after December 15, 2016; however, earlier application is permitted. We elected to early adopt Update No. 2015-17 on a prospective basis; as such, prior periods were not retrospectively adjusted. The adoption of Update No. 2015-17 did not have a material impact on our financial position or results of operations.

ASC Update No. 2016-07

In March 2016, the FASB issued ASC Update No. 2016-07, Investments - Equity Method and Joint Ventures (Topic 323). When a previously held investment qualifies for equity method accounting due to an increase in ownership

interest or influence, Update 2016-07 eliminates the requirement for investors to adjust results retroactively as if the equity method had been in effect during prior periods the investment was held. Instead, it requires investors to adopt the equity method of accounting as of the date the investment becomes qualified for equity method accounting. We elected to early adopt Update No. 2016-07 on a prospective basis.
The adoption of Update No. 2016-07 did not impact on our financial position or results of operations.

Standards to be Implemented

ASC Updates No. 2014-09, No. 2016-08, No. 2016-10, No. 2016-11 and No. 2016-12

In May 2014, the FASB issued ASC Update No. 2014-09, Revenue from Contracts with Customers (Topic 606). Update No. 2014-09 provides enhancements to the quality and consistency of how revenue is reported while also improving comparability in the financial statements of companies using International Financial Reporting Standards and U.S. GAAP. The core principle requires entities to recognize revenue in a manner that depicts the transfer of goods or services to customers in amounts that reflect

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the consideration an entity expects to be entitled to in exchange for those goods or services. In July 2015, the FASB voted to approve a one year deferral, making the standard effective for public entities for annual and interim periods beginning after December 15, 2017.

In March 2016, the FASB issued ASC Update No. 2016-08, Revenue from Contracts with Customers (Topic 606): Principal versus Agent Considerations (Reporting Revenue Gross versus Net). The purpose of Update No. 2016-08 is to clarify the guidance on principal versus agent considerations. It includes indicators that help to determine whether an entity controls the specified good or service before it is transferred to the customer and to assist in determining when the entity satisfied the performance obligation and as such, whether to recognize a gross or a net amount of consideration in their consolidated statement of operations.

In April 2016, the FASB issued ASC Update No. 2016-10, Revenue from Contracts with Customers (Topic 606): Identifying Performance Obligations and Licensing. Update No. 2016-10 clarifies that entities are not required to assess whether promised goods or services are performance obligations if they are immaterial in the context of the contract. Update 2016-10 also addresses how to determine whether promised goods or services are separately identifiable and permits entities to make a policy election to treat shipping and handling costs as fulfillment activities. In addition, it clarifies key provisions in Topic 606 related to licensing.

In May 2016, the FASB issued ASC Update No. 2016-11, Revenue Recognition (Topic 605) and Derivatives and Hedging (Topic 815). Update No. 2016-11 rescinds previous SEC comments that were codified in Topic 605, Topic 932 and Topic 815. Upon adoption of ASC 606, certain SEC comments including guidance on accounting for shipping and handling fees and costs and consideration given by a vendor to a customer should not be relied upon.

In May 2016, the FASB also issued ASC Update No. 2016-12, Revenue from Contracts with Customers (Topic 606): Narrow Scope Improvements and Practical Expedients. Update No. 2016-12 provides clarity around collectability, presentation of sales taxes, non-cash consideration, contract modifications at transition and completed contracts at transition. Update No. 2016-12 also includes a technical correction within ASC 606 related to required disclosures if the guidance is applied retrospectively upon adoption.

We expect to adopt Topic 606, and the aforementioned updates, effective January 1, 2018. We are in the process of determining the effect that the adoption of these standards will have on our financial position and results of operations.

ASC Update No. 2016-01

In January 2016, the FASB issued ASC Update No. 2016-01, Financial Instruments-Overall (Subtopic 825-10): Recognition and Measurement of Financial Assets and Financial Liabilities. It is effective for fiscal years beginning after December 15, 2017, including interim periods within those fiscal years. Early application of certain provisions is permitted. Update 2016-01 requires entities to measure equity investments that do not result in consolidation and are not accounted for under the equity method at fair value with changes recognized in net income. However, an entity may choose to measure equity investments that do not have readily determinable fair values at cost minus impairment, if any, plus or minus changes resulting from observable price changes in orderly transactions for the identical or a similar investment of the same issuer. It also simplifies the impairment assessment of equity investments without readily determinable fair values by requiring a qualitative assessment to identify impairment. Update 2016-01 also requires separate presentation of financial assets and financial liabilities by measurement category and form of financial asset and liability. The adoption of Update No. 2016-01 is not expected to have a material impact on our financial position or results of operations.

ASC Update No. 2016-02

In February 2016, the FASB issued ASC Update No. 2016-02, Leases (Topic 842). It is effective for fiscal years beginning after December 15, 2018, including interim periods within those fiscal years. Earlier application is permitted. Update 2016-02 is intended to increase the transparency and comparability among organizations by recognizing lease asset and lease liabilities on the balance sheet, including those previously classified as operating leases under current GAAP, and disclosing key information about leasing arrangements. We are in the process of determining the effect that the adoption of this standard will have on our financial position and results of operations.

ASC Update No. 2016-09

In March 2016, the FASB issued ASC Update No. 2016-09, Compensation- Stock Compensation (Topic 718): Improvements to Employee Share-Based Payment Accounting. This update is effective for annual reporting periods after December 15, 2016, including interim periods within those fiscal periods. Early adoption is permitted. The purpose of the update is to simplify several areas of the accounting for share-based payment transactions, including the income tax consequences, classification of awards as

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either equity or liabilities, and classification on the statement of cash flows. We are in the process of determining the effect that the adoption will have on our financial position and results of operations.

ASC Update No. 2016-13

In June 2016, the FASB issued ASC Update No. 2016-13, Financial Instruments - Credit Losses (Topic 326): Measurement of Credit Losses on Financial Instruments. The update is effective for fiscal years beginning after December 15, 2019. Early adoption is permitted for fiscal years beginning after December 15, 2018. The purpose of Update No. 2016-13 is to replace the current incurred loss impairment methodology for financial assets measured at amortized cost with a methodology that reflects expected credit losses and requires consideration of a broader range of reasonable and supportable information, including forecasted information, to develop credit loss estimates. We are in the process of determining the effect that the adoption will have on our financial position and results of operations.

No other new accounting pronouncements, issued or effective, during the period had, or is expected to have, a material impact on our condensed consolidated financial statements.

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

Introduction

Boston Scientific Corporation is a worldwide developer, manufacturer and marketer of medical devices that are used in a broad range of interventional medical specialties. Our mission is to transform lives through innovative medical solutions that improve the health of patients around the world. Our products and technologies are used to diagnose or treat a wide range of medical conditions, including heart, vascular, digestive, pulmonary, urological, pelvic health, and chronic pain conditions. We continue to innovate in these areas and are intent on extending our innovations into new geographies and high-growth adjacency markets.

AMS Portfolio Acquisition

On August 3, 2015, we completed the acquisition of the American Medical Systems male urology portfolio (AMS Portfolio Acquisition), which includes the men's health and prostate health businesses, from Endo International plc. Total consideration was comprised of $1.616 billion in up-front cash plus related fees and expenses, and a potential additional $50 million in consideration based on 2016 sales. The AMS male urology portfolio is being integrated with our formerly named Urology and Women's Health business, and the joint businesses have become Urology and Pelvic Health.

Financial Summary

Three Months Ended June 30, 2016

Our net sales for the second quarter of 2016 were $2.126 billion, as compared to net sales of $1.843 billion for the second quarter of 2015, an increase of $283 million, or 15 percent. Our adjusted net sales, which excludes a negative impact of $24 million in the second quarter 2016, due to changes in foreign currency exchange rates, increased $307 million, or 16 percent, as compared to the same period in the prior year.[1] This increase included adjusted net sales of approximately $102 million in the second quarter 2016, with no prior year period related net sales, due to the acquisition of the AMS Portfolio Acquisition, which included the men's health and prostate health businesses, from Endo International plc, during the third quarter of 2015. Refer to Quarterly Results and Business Overview for a discussion of our net sales by global business.

Our reported net loss for the second quarter of 2016 was $207 million, or $(0.15) per share, due to litigation-related charges. Our reported results for the second quarter of 2016 included acquisition- and divestiture-related net charges, restructuring and restructuring-related net charges, litigation-related net charges and amortization expense totaling $580 million (after-tax), or $0.42 per share. Adjusted net income, which excludes these items, for the second quarter of 2016 was $373 million, or $0.27 per share.[1] Our reported net income for the second quarter of 2015 was $102 million, or $0.08 per share. Our reported results for the second quarter of 2015 included an intangible asset impairment charge, acquisition- and divestiture-related net charges, restructuring and restructuring-related net charges, litigation-related net credits, debt extinguishment charges and amortization expense totaling $192 million (after-tax), or $0.14 per share. Adjusted net income, which excludes these items, for the second quarter of 2015 was $294 million, or $0.22 per share.[1]

[1] Adjusted net sales growth rates, which exclude the impact of changes in foreign currency exchange rates, and adjusted net income and adjusted net income per share, which exclude certain items required by GAAP, are not prepared in accordance with U.S. GAAP. Refer to Additional Information for a discussion of management's use of these non-GAAP financial measures.

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The following is a reconciliation of our results of operations prepared in accordance with U.S. GAAP to those adjusted results considered by management. Refer to Quarterly Results and Business Overview for a discussion of each reconciling item:

	Three Months Ended June 30, 2016			
in millions, except per share data	Pre-Tax	Tax Impact	After-Tax	Impact per share
GAAP net income (loss)	$(397)	$ 190	$ (207)	$(0.15)
Non-GAAP adjustments:				
Acquisition- and divestiture-related net charges	53	(4)	49	0.04 *
Restructuring and restructuring-related net charges	26	(5)	21	0.02 *
Litigation-related net charges	618	(224)	394	0.28 *
Amortization expense	135	(19)	116	0.08 *
Adjusted net income	$435	$ (62)	$ 373	$0.27

*Assumes dilution of 17.7 million shares for the three months ended June 30, 2016 for all or a portion of these non-GAAP adjustments.

	Three Months Ended June 30, 2015			
in millions, except per share data	Pre-Tax	Tax Impact	After-Tax	Impact per share
GAAP net income (loss)	$105	$ (3)	$ 102	$0.08
Non-GAAP adjustments:				
Intangible asset impairment charge	9	(2)	7	0.01
Acquisition- and divestiture-related net charges	49	(7)	42	0.03
Restructuring and restructuring-related net charges	16	(2)	14	0.01
Litigation-related net credits	(1)	—	(1)	0.00
Debt extinguishment charges	45	(16)	29	0.02
Amortization expense	116	(15)	101	0.07
Adjusted net income	$339	$ (45)	$ 294	$0.22

Cash provided by operating activities was $422 million in the second quarter of 2016, as compared to $60 million in the second quarter of 2015. As of June 30, 2016, we had total debt of $5.427 billion, cash and cash equivalents of $438 million and working capital of $148 million. Refer to Liquidity and Capital Resources for further discussion.

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Six Months Ended June 30, 2016

Our net sales for the first half of 2016 were $4.090 billion, as compared to net sales of $3.611 billion for the first half of 2015, an increase of $479 million, or 13 percent. Our adjusted net sales, which excludes a negative impact of $82 million on our first half of 2016 net sales, due to changes in foreign currency exchange rates, increased $561 million, or 15 percent, as compared to the same period in the prior year.[1] This increase included adjusted net sales of approximately $200 million in the first half of 2016, with no prior year period related net sales, due to the acquisition of the AMS Portfolio Acquisition, as previously described. Refer to Quarterly Results and Business Overview for a discussion of our net sales by global business.

Our reported net loss for the first half of 2016 was $5 million, or $(0.00) per share. Our reported results for the first half of 2016 included acquisition- and divestiture-related net charges, restructuring and restructuring-related net charges, litigation-related net charges, and amortization expense totaling $756 million (after-tax), or $0.55 per share. Excluding these items, net income for the first half of 2016 was $751 million, or $0.55 per share.[1] Our reported net income for the first half of 2015 was $101 million, or $0.07 per share. Our reported results for the first half of 2015 included intangible asset impairment charges, acquisition- and divestiture-related net charges, restructuring and restructuring-related net charges, litigation-related net charges, debt extinguishment charges, pension termination charges and amortization expense totaling $479 million (after-tax), or $0.36 per share. Excluding these items, net income for the first half of 2015 was $580 million, or $0.43 per share.[1]

[1] Sales growth rates that exclude the impact of sales from divested businesses and/or changes in foreign currency exchange rates and net income and net income per share excluding certain items required by GAAP are not prepared in accordance with U.S. GAAP. Refer to Additional Information for a discussion of management's use of these non-GAAP financial measures.

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The following is a reconciliation of our results of operations prepared in accordance with U.S. GAAP to those adjusted results considered by management. Refer to Quarterly Results and Business Overview for a discussion of each reconciling item:

	Six Months Ended June 30, 2016				
in millions, except per share data	Pre-Tax	Tax Impact	After-Tax	Impact per share	
GAAP net income (loss)	$(169)	$164	$ (5)	$(0.00)	
Non-GAAP adjustments:					
Acquisition- and divestiture-related net charges	96	(2)	94	0.07	*
Restructuring and restructuring-related net charges	38	(10)	28	0.02	*
Litigation-related net charges	628	(228)	400	0.29	*
Amortization expense	271	(37)	234	0.17	*
Adjusted net income	$864	$(113)	$ 751	$0.55	

*Assumes dilution of 18.6 million shares for the six months ended June 30, 2016 for all or a portion of these non-GAAP adjustments.

	Six Months Ended June 30, 2015			
in millions, except per share data	Pre-Tax	Tax Impact	After-Tax	Impact per share
GAAP net income (loss)	$54	$47	$ 101	$0.07
Non-GAAP adjustments:				
Intangible asset impairment charges	9	(2)	7	0.01
Acquisition- and divestiture-related net charges	91	(5)	86	0.07
Restructuring and restructuring-related net charges	37	(6)	31	0.02
Litigation-related net charges	192	(70)	122	0.09
Debt extinguishment charges	45	(16)	29	0.02
Pension termination charges	8	(3)	5	0.00
Amortization expense	229	(30)	199	0.15
Adjusted net income	$665	$(85)	$ 580	$0.43

Cash provided by operating activities was $537 million in the first half of 2016, as compared to cash used for operating activities of $137 million in the first half of 2015. The increase was primarily due to litigation-related payments of $600 million during the first half of 2015 that did not recur in the first half of 2016.

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Quarterly Results and Business Overview

Net Sales

The following table provides our net sales by business and the relative change on an as reported and constant currency basis. The constant currency growth rates in the tables below can be recalculated from our net sales presented in Note K – Segment Reporting to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q. Constant currency growth rates, which exclude the impact of changes in foreign currency exchange rates, are not financial measures prepared in accordance with U.S. GAAP and should not be considered in isolation from, or as a replacement for, the most directly comparable GAAP financial measure. Refer to Additional Information for a further discussion of management's use of this non-GAAP financial measure.

(in millions)	Three Months Ended June 30, 2016	2015	Change As Reported Currency Basis	Less: Impact of Foreign Currency	Constant Currency Basis
Interventional Cardiology	$579	$515	13%	1 %	12 %
Peripheral Interventions	258	228	13%	(1) %	14 %
Cardiovascular	837	743	13%	0 %	13 %
Cardiac Rhythm Management	477	460	4 %	0 %	4 %
Electrophysiology	60	57	5 %	(1) %	6 %
Rhythm Management	537	517	4 %	(1) %	5 %
Endoscopy	361	326	11%	0 %	11 %
Urology and Pelvic Health	256	135	89%	(1) %	90 %
Neuromodulation	135	122	11%	(1) %	12 %
MedSurg	752	583	29%	0 %	29 %
Net Sales	$2,126	$1,843	15%	(1) %	16 %

Growth rates are based on actual, non-rounded amounts and may not recalculate precisely.

(in millions)	Six Months Ended June 30, 2016	2015	Change As Reported Currency Basis	Less: Impact of Foreign Currency	Constant Currency Basis
Interventional Cardiology	$1,128	$1,010	12 %	(1) %	13 %
Peripheral Interventions	501	445	12 %	(2) %	14 %
Cardiovascular	1,629	1,455	12 %	(1) %	13 %
Cardiac Rhythm Management	910	916	(1)%	(2) %	1 %
Electrophysiology	119	115	4 %	(2) %	6 %
Rhythm Management	1,029	1,031	(0)%	(1) %	1 %
Endoscopy	693	631	10 %	(1) %	11 %
Urology and Pelvic Health	483	258	87 %	(2) %	89 %
Neuromodulation	256	236	8 %	(2) %	10 %
MedSurg	1,432	1,125	27 %	(1) %	28 %

Net Sales	$4,090	$3,611	13 %	(2) %	15 %

Growth rates are based on actual, non-rounded amounts and may not recalculate precisely.

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Cardiovascular

Interventional Cardiology

Our Interventional Cardiology business develops, manufactures and markets technologies for diagnosing and treating coronary artery disease and other cardiovascular disorders. Product offerings include coronary stents, including drug-eluting and bare metal stent systems, balloon catheters, rotational atherectomy systems, guide wires, guide catheters, embolic protection devices, crossing and re-entry devices for the treatment of chronically occluded coronary vessels, diagnostic catheters used in percutaneous transluminal coronary angioplasty procedures, and intravascular ultrasound (IVUS) imaging systems. We also offer structural heart products in certain markets, which include a device for transcatheter aortic valve replacement (TAVR) and a device designed to close the left atrial appendage in patients with atrial fibrillation that are at risk for ischemic stroke.

Our drug-eluting stent systems include our next generation SYNERGY™ Everolimus-Eluting Platinum Chromium Coronary Stent System and our Promus PREMIER™ Everolimus-Eluting Platinum Chromium Coronary Stent System, both of which are designed to provide physicians with improved drug-eluting stent performance in treating patients with coronary artery disease. SYNERGY™ features an ultra-thin abluminal (outer) bioabsorbable polymer coating, while PREMIER™ features a unique customized platinum chromium alloy stent architecture and an enhanced stent delivery system. We received FDA and Japanese regulatory approval of the SYNERGY™ technology in the fourth quarter of 2015 and we launched SYNERGY™ in Japan in the first quarter of 2016.

Our structural heart product offerings include our Lotus™ Valve System, a device for TAVR, and our WATCHMAN® device designed to close the left atrial appendage in patients with non-valvular atrial fibrillation who are at risk for ischemic stroke. The Lotus™ Valve System consists of a stent-mounted tissue valve prosthesis and catheter delivery system for guidance and placement of the valve. The Lotus™ Valve System is CE-marked in the European Union (EU), and in the U.S., it is an investigational device and not available for sale. At the end of 2015, we completed enrollment in our REPRISE III clinical trial and expect FDA approval of the Lotus Valve System in late 2017. The WATCHMAN® Left Atrial Appendage Closure Technology (WATCHMAN®) is the first device studied in a randomized clinical trial to offer an alternative to warfarin, and is marketed in CE-mark countries and other international countries, as well as the U.S. following FDA approval in March 2015. We believe that Watchman® will be the only LAAC technology commercially available in the U.S. for multiple years. In November 2015, we received CE Mark for our next generation device, Watchman FLX™. Shortly after approval, we began a European initial market release of Watchman FLX. The initial market release was suspended near the end of the first quarter of 2016 due to a higher than expected rate of device embolization. Following an extensive data evaluation, we have decided to pursue potential design enhancements prior to returning a next generation device to market.

Our net sales of Interventional Cardiology products of $579 million represented 27 percent of our consolidated net sales for the second quarter of 2016. Our Interventional Cardiology net sales increased $64 million, or 13 percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $4 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $68 million, or 12 percent, as compared to the same period in the prior year. This year-over-year increase was primarily related to sales of our drug-eluting stents, led by our ongoing global launch of the SYNERGY™ stent, our WATCHMAN® device following the U.S. commercial launch during the first quarter of 2015, and our Lotus™ Valve System in the EU; along with operational growth in our other cardiology product lines, including our PCI Guidance System product offerings and our portfolio of products for treating complex cardiovascular disease.

Peripheral Interventions

Our Peripheral Interventions (PI) product offerings include stents, balloon catheters, wires, peripheral embolization devices and other devices used to diagnose and treat peripheral vascular disease, along with certain products to treat, diagnose and ease various forms of cancer.

Our net sales of PI products of $258 million represented 12 percent of our consolidated net sales for the second quarter of 2016. Our PI net sales increased $30 million, or 13 percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $3 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $33 million, or 14 percent, as compared to the same period in the prior year. This year-over-year increase was primarily driven by revenues from our Atherectomy and Thrombectomy systems, as well as growth in our core PI franchises, particularly our stent franchise following FDA approval and launch of our Innova™ Vascular self-expanding stent system in the U.S. and Japan, our interventional oncology franchise and our drug-eluting product franchise.

On December 31, 2015, we acquired the interventional radiology portfolio of CeloNova Biosciences (CeloNova). The acquisition includes drug-eluting microspheres designed to be loaded with chemotherapy drugs for delivery to cancerous tumors, and spherical

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embolic products used to treat uterine fibroids and other conditions. We believe the CeloNova team and technologies will help advance our position and growth profile within the interventional oncology market.

Rhythm Management

Cardiac Rhythm Management

Our Cardiac Rhythm Management (CRM) business develops, manufactures and markets a variety of implantable devices including implantable cardioverter defibrillator (ICD) systems and implantable cardiac resynchronization therapy defibrillators, including the world's first and only commercially available subcutaneous implantable cardioverter defibrillator, the S-ICD® System, and pacemaker systems that monitor the heart and deliver electricity to treat cardiac abnormalities. In addition, in most geographies, our implantable device systems include our remote LATITUDE® Patient Management System, which enables physicians to monitor device performance remotely, allowing for more frequent monitoring in order to guide treatment decisions.

Our net sales of CRM products of $477 million represented 23 percent of our consolidated net sales for the second quarter of 2016. Our net sales of CRM products increased $17 million, or four percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $5 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $22 million, or four percent, as compared to the same period in the prior year. This year-over-year increase was primarily driven by strong global pacemaker growth including the launch of our ACCOLADE™ family of magnetic resonance imaging (MRI) safe pacemakers and the Ingevity™ MRI pacing lead in the U.S., global growth from our quadripolar cardiac resynchronization therapy pacemakers (CRT-P), strong Japan CRM sales with S-ICD and our MRI safe technologies, and benefits from our sales collaboration agreement with Preventice, Inc. In the U.S., the second quarter of 2016 was the first full quarter of commercialization for our Acuity X4 quadripolar LV pacing lead in both the cardiac resynchronization therapy defibrillator (CRT-D) and CRT-P franchises. This launch helped offset lower volumes of replacement procedures for our defibrillators due to their extended longevity and pressure from high voltage MRI technologies primarily in the U.S.

The following are the components of our CRM net sales:

	Three Months Ended June 30,	
(in millions)	2016	2015
Defibrillator systems	$333	$335
Pacemaker systems	144	125
CRM products	$477	$460

We market several lines of ICD’s, including our line of MINIs, the world’s smallest, thinnest ICD, and our line of ELs (extended longevity), the world’s longest lasting ICD due to our proprietary EnduraLife™ battery technology. In addition, we offer our EMBLEM™ S-ICD system, which affords physicians the ability to treat patients who are at risk for sudden cardiac arrest without touching the heart or invading the vasculature. Our EMBLEM™ S-ICD system offers greater longevity, LATITUDE® Patient Management remote monitoring technology and smaller size as compared to the prior generation. In April 2016, we received CE Mark approval for the new EMBLEM™ MRI S-ICD System, as well as magnetic resonance conditional labeling for all previously implanted EMBLEM S-ICD Systems, and began commercialization of our next generations S-ICD system in Europe. We also offer several lines of CRT-D systems, including our X4 line of quadripolar systems and Acuity X4 quadripolar LV leads, and the ACUITY™ PRO lead delivery system. We initiated the full launch of our X4 quadripolar CRT-D systems in Japan and Australia in the first

quarter of 2015 and in February of 2016 we received FDA approval for the Acuity™ X4 Quadripolar lead.

We market our ACCOLADE™ family of pacemaker systems in the U.S., Europe, and Japan. We received FDA approval of our ACCOLADE™ MRI-compatible pacemaker in April 2016 and we expect FDA approval of our EMBLEM MRI-compatible system in third quarter of 2016. Our cardiac resynchronization therapy pacemaker product offerings include our newest generation VISIONIST™ and VALITUDE X4 quadripolar CRT-P devices, which are built on the same platform as our high voltage cardiac resynchronization therapy defibrillator, are enabled for remote patient monitoring, and include features that promote ease of use.

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Electrophysiology

Our Electrophysiology business develops less-invasive medical technologies used in the diagnosis and treatment of rate and rhythm disorders of the heart. Our leading products include the Blazer™ line of ablation catheters, designed to deliver enhanced performance and responsiveness, and the Rhythmia™ Mapping System, a next-generation, catheter-based, 3-D cardiac mapping and navigation solution designed to help diagnose and treat a variety of arrhythmias.

Our net sales of Electrophysiology products of $60 million represented three percent of our consolidated net sales for the second quarter of 2016. Our Electrophysiology net sales increased $3 million, or five percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $1 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $4 million, or six percent, as compared to the same period in the prior year. This year-over-year increase was primarily driven by increased sales of our Rhythmia™ Mapping System and related products. In the first quarter of 2016, we initiated a full European launch of our Blazer IntellaNav™ OI catheter which is used with our Rhythmia™ Mapping System and, in July of 2016, we received FDA approval for this same catheter. In the second quarter of 2016, we received FDA approval for IntellaNav™ XP and the IntellaNav MiFi™ XP navigation-enabled ablation catheters that are used with the Rhythmia™ Mapping System. The second quarter of 2016 also marked the start of commercializing for our next generation IntellaTip™ MiFi OI catheter in select international markets. Finally, we received FDA approval for our Blazer™ Open Irrigated System with Atrial Flutter indication with full U.S. commercialization beginning in the second quarter of 2016.

MedSurg

Endoscopy

Our Endoscopy business develops and manufactures devices to treat a variety of medical conditions including diseases of the digestive and pulmonary space.

Our net sales of Endoscopy products of $361 million represented 17 percent of our consolidated net sales for the second quarter of 2016. Our Endoscopy net sales increased $35 million, or 11 percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $3 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $38 million, or 11 percent, as compared to the same period in the prior year. This year-over-year increase was primarily driven by growth across several of our key product franchises, including our biliary device franchise with the launch of SpyGlass™ DS Direct Visualization System and our AXIOS Stent and Electrocautery-Enhanced Delivery System for endoscopic ultrasound-guided transmural drainage of pancreatic pseudocysts; our metal stent franchise driven by our Biliary WallFlex® product family; and our hemostasis franchise, featuring our Resolution™ and Resolution 360™ Clips.

On April 2, 2015, we acquired Xlumena, Inc. (Xlumena), a medical device company that developed minimally invasive devices for Endoscopic Ultrasound (EUS) guided transluminal drainage of targeted areas within the gastrointestinal tract.

Urology and Pelvic Health

Our Urology and Pelvic Health business develops and manufactures devices to treat various urological and pelvic conditions. Our net sales of Urology and Pelvic Health products of $256 million represented 12 percent of our consolidated net sales for the second quarter of 2016. Urology and Pelvic Health net sales increased $121 million, or 89 percent, in the second quarter of 2016, as compared to the same period in the prior year. Our adjusted net sales,

which excludes a negative impact of $7 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $128 million, or 90 percent, as compared to the same period in the prior year. This year-over-year increase was primarily attributable to revenue of approximately $102 million from sales of products acquired with the AMS Portfolio Acquisition along with growth across all of our other global franchises, including our Pelvic Floor franchise as a result of market share gains primarily driven by a competitor exiting the market during the first quarter of 2016.

On August 3, 2015, we completed the AMS Portfolio Acquisition, which included the men's health and prostate health businesses, from Endo International plc., for $1.616 billion in up-front cash plus related fees and expenses, and a potential additional $50 million in consideration based on 2016 sales. The AMS Portfolio Acquisition includes the procurement of leading products for the treatment of a variety of urologic conditions, including the minimally invasive GreenLight XPS™ and HPS™ Laser Therapy Systems for treating benign prostatic hyperplasia, the AMS 700™ Inflatable Penile Prosthesis for treating erectile dysfunction, and the AMS 800™ Urinary Control System for treating male stress urinary incontinence.

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Neuromodulation

Our Neuromodulation business offers the Precision™, Precision Spectra™ and Precision Novi™Spinal Cord Stimulator (SCS) Systems, used for the management of chronic pain. The Precision Spectra System is the world's first and only SCS system with 32 contacts and 32 dedicated power sources and is designed to provide improved pain relief to a wide range of patients who suffer from chronic pain. The Precision Novi™ SCS System offers patients and physicians with the smallest 16-contact high capacity primary cell (PC), also referred to as non-rechargeable, device for the treatment of chronic pain. In May 2016, we launched the Precision Montage™ MRI SCS System after receiving FDA approval. The Precision Montage System offers customized relief to patients with chronic pain while also enabling safe access to full-body MRI technology. We have CE mark approval for Vercise™ Deep Brain Stimulation (DBS) System in Europe for the treatment of Parkinson's disease, tremor and intractable primary and secondary dystonia, a neurological movement disorder characterized by involuntary muscle contractions. In September 2015, we gained CE mark approval for the Vercise™ PC DBS System with its Neural Navigator™ programming software and Vercise Cartesia™ Directional Lead. The system allows for programming flexibility, designed to treat a greater range of patients throughout their disease progression. The Cartesia™ Directional Lead uses multi-directional stimulation for greater precision, intended to minimize side effects for patients. We are currently in U.S. pivotal trial with our Vercise™ DBS System for the treatment of Parkinson’s disease.

Our net sales of Neuromodulation products of $135 million represented six percent of our consolidated net sales for the second quarter of 2016. Our Neuromodulation net sales increased $13 million, or 11 percent, in the second quarter of 2016, as compared the same period in the prior year. Our adjusted net sales, which excludes a negative impact of $1 million in the second quarter of 2016, due to changes in foreign currency exchange rates, increased $14 million, or 12 percent, as compared to the same period in the prior year. This year-over-year increase was primarily driven by share gains from our Montage™ System, continued adoption of the Precision Spectra™ SCS System in the U.S. and increased net sales in Europe, driven by our Vercise™ DBS Systems and non-rechargeable Precision Novi™ SCS System.

On July 27, 2016, we acquired Cosman Medical, Inc. (Cosman), a privately held manufacturer of radiofrequency ablation systems, expanding our Neuromodulation portfolio and offering physicians treating patients with chronic pain a wider choice of non-opioid therapeutic options.

Emerging Markets

As part of our strategic imperatives to drive global expansion, described in our most recent Annual Report on Form 10-K, we are seeking to grow net sales and market share by expanding our global presence, including in Emerging Markets. We define Emerging Markets as 20 countries that we believe have strong growth potential based on their economic conditions, healthcare sectors, and our global capabilities. We are seeking to expand our presence and strengthen relationships in order to grow net sales and market share within our Emerging Markets, and we have increased our investment in infrastructure in these countries in order to maximize opportunities. Our Emerging Markets net sales represented approximately nine percent of our consolidated net sales in the second quarter of 2016 and 10 percent of our consolidated net sales in the second quarter of 2015; however, in the second quarter of 2016, these sales grew eight percent compared to the prior year period.

Gross Profit

Our gross profit was $1.487 billion for the second quarter of 2016, $1.303 billion for the second quarter of 2015, $2.879 billion for the first half of 2016 and $2.551 billion for the first half of 2015. As a percentage of net sales, our gross profit decreased to 70.0 percent in the second quarter of 2016, as compared to 70.7 percent in the second quarter of 2015 and decreased to 70.4 percent in the first half of 2016, as compared to 70.6 percent in the first half of 2015. The following is a reconciliation of our gross profit margins and a description of the drivers of the change from period

to period:

	Three Months	Six Months
Gross profit margin - period ended June 30, 2015	70.7 %	70.6 %
Manufacturing cost reductions	2.1	2.0
Sales pricing and mix	(0.1)	(0.1)
Net impact of foreign currency	(1.2)	(0.7)
All other, including other period expense	(1.5)	(1.4)
Gross profit margin - period ended June 30, 2016	70.0 %	70.4 %

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The primary factor contributing to the decrease in our gross profit margin during the second quarter and first half of 2016, as compared to the same periods in 2015, was net negative impacts of foreign currency fluctuations and other period expense partially offset by the positive impact of cost reductions resulting from our restructuring and other process improvement programs.

Operating Expenses

The following table provides a summary of certain of our operating expenses:

	Three Months Ended June 30,				Six Months Ended June 30,			
	2016		2015		2016		2015	
(in millions)	$	% of Net Sales	$	% of Net Sales	$	% of Net Sales	$	% of Net Sales
Selling, general and administrative expenses	779	36.6%	700	38.0%	1,497	36.6%	1,367	37.9%
Research and development expenses	222	10.4%	220	11.9%	431	10.5%	412	11.4%
Royalty expense	20	0.9 %	18	1.0 %	39	1.0 %	36	1.0 %

Selling, General and Administrative (SG&A) Expenses

In the second quarter of 2016, our SG&A expenses increased $79 million, or 11 percent, as compared to the second quarter of 2015, and were 140 basis points lower as a percentage of net sales. In the first half of 2016, our SG&A expenses increased $130 million, or 10 percent, as compared to the first half of 2015, but were 130 basis points lower as a percentage of net sales. The decrease in SG&A as a percentage of sales was primarily driven by the recent suspension of the medical device tax, as well as our targeted initiatives that have been focused on reducing SG&A.

Research and Development (R&D) Expenses

In the second quarter of 2016, our R&D expenses increased $2 million, or one percent, as compared to the second quarter of 2015, but were 150 basis points lower as a percentage of net sales. In the first half of 2016, our R&D expenses increased $19 million, or five percent, as compared to the first half of 2015, but were 90 basis points lower as a percentage of net sales. We remain committed to advancing medical technologies and investing in meaningful research and development projects across our businesses in order to maintain a healthy pipeline of new products that we believe will contribute to profitable sales growth and increased costs related to recent acquisitions and alliances.

Royalty Expense

In the second quarter of 2016, our royalty expense increased $2 million, or 11 percent, as compared to the second quarter of 2015. In the first half of 2016, our royalty expense increased $3 million, or eight percent, as compared to the first half of 2015. Although our royalty expense increased period over period, it remained relatively flat at approximately one percent of net sales in the both second quarters and first halves of 2016 and 2015. The consistency as a percentage of net sales year-over-year relates primarily to a renegotiation of a royalty agreement in 2014 that resulted in a lower royalty rate structure.

Amortization Expense

Our amortization expense was $135 million in the second quarter of 2016, as compared to $116 million in the second quarter of 2015, and $271 million in the first half of 2016, as compared to $229 million in the first half of 2015. This increase was primarily due to amortizable intangible assets acquired as part of the AMS portfolio acquisition during

the third quarter of 2015. Amortization expense is excluded by management for purposes of evaluating operating performance.

Intangible Asset Impairment Charges

We incurred intangible asset impairment charges, including charges for impairments of in-process research and development, of $9 million during the second quarter and first half of 2015. No intangible asset impairment chargers were recorded in the first half of 2016 .

Refer to Note C - Goodwill and Other Intangible Assets to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for additional details related to our intangible asset impairment charges. Intangible asset impairment charges are excluded by management for purposes of evaluating operating performance.

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Contingent Consideration Expense

We recorded net expenses of $33 million and $37 million during the second quarter and first half of 2016, respectively, and net expenses of $19 million and $46 million during the second quarter and first half of 2015, respectively, related to the change in fair value of our contingent consideration liabilities. Refer to Note B - Acquisitions and Strategic Investments to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for additional details related to our contingent consideration expenses. Contingent consideration expense is excluded by management for purposes of evaluating operating performance.

Restructuring Charges and Restructuring-related Activities

We have two active restructuring programs, our 2016 Restructuring Plan, which was approved on June 6, 2016 and commenced activities at the end of the second quarter of 2016, and our 2014 Restructuring Plan, which was approved on October 22, 2013 and substantially completed at the end of 2015, with the exception of certain actions associated with our Plant Network Optimization strategy. Our 2016 Restructuring Plan is designed to remain active through the end of 2018.

We estimate that the 2016 Restructuring Plan will reduce our gross annual expenses by approximately $115 million to $150 million by the end of 2020 as program benefits are realized, and expect a portion of the Program savings to be reinvested in strategic growth initiatives. We estimate the implementation of the 2016 Restructuring Plan will result in total pre-tax charges of approximately $175 million to $225 million, of which approximately $160 million to $210 million is expected to result in cash outlays. We have recorded related costs of $19 million since the inception of the 2016 Restructuring Plan.

We estimate that the 2014 Restructuring Plan will reduce our gross annual expenses by approximately $200 million by the end of 2016, and we expect a substantial portion of the savings to be reinvested in growth initiatives. We estimate that the implementation of the 2014 Restructuring Plan will result in total pre-tax charges of approximately $255 million to $270 million, of which approximately $240 million to $255 million is expected to result in cash outlays. We have recorded costs of $249 million since the inception of the 2014 Restructuring Plan.

We recorded restructuring charges pursuant to our restructuring plans of $14 million in the second quarter of 2016, $3 million in the second quarter of 2015, $17 million during the first half of 2016 and $9 million during the first half of 2015. In addition, we recorded expenses within other lines of our accompanying unaudited condensed consolidated statements of operations related to our restructuring initiatives of $12 million in the second quarter of 2016, $12 million in the second quarter of 2015, $22 million in the first half of 2016 and $28 million in the first half of 2015. Restructuring and restructuring-related costs are excluded by management for purposes of evaluating operating performance.

We made cash payments of $40 million during the first half of 2016 and $46 million during the first half of 2015, associated with our restructuring initiatives.

Refer to Note F - Restructuring Related Activities to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for additional details related to our restructuring plans.

Litigation-related charges and credits

We recorded litigation-related net charges of $618 million in the second quarter of 2016 and $628 million in the first half of 2016. We recorded litigation-related net credits of $1 million in the second quarter of 2015 and

litigation-related net charges of $192 million in the first half of 2015. Litigation-related charges and credits are excluded by management for purposes of evaluating operating performance. Refer to Note I – Commitments and Contingencies to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for discussion of our material legal proceedings.

Pension termination charges

We recorded pension termination charges of $8 million during the first quarter of 2015 and an additional $36 million during the third quarter of 2015 for a total of $44 million of pension termination charges in the year ended December 31, 2015. These charges were associated with the termination of the Guidant Retirement Plan, a frozen defined benefit plan. No pension termination charges were recorded during the first half of 2016, and we do not expect to incur any additional charges in the future related to the termination of the Guidant Retirement Plan. The pension termination charges are excluded by management for purposes of evaluating operating performance.

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Interest Expense

Our interest expense was $59 million in the second quarter of 2016 and $118 million during the first half of 2016, as compared to $106 million in the second quarter of 2015 and $167 million during the first half of 2015. Our average borrowing rate was 4.0 percent in the second quarter of 2016 and 4.0 percent in the first half of 2016 and 8.0 percent the second quarter of 2015 and 6.9 percent in the first half of 2015. Refer to Liquidity and Capital Resources and Note D - Fair Value Measurements and Note E – Borrowings and Credit Arrangements to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for information regarding our debt obligations and related derivative instruments and hedging activities.

Other, net

Our other, net reflected expense of $4 million and $8 million in the second quarter of 2016 and 2015, respectively. During the first half of 2016 and 2015, our other, net reflected expense of $10 million and expense of $22 million, respectively. The following are the components of other, net:

	Three Months Ended June 30,		Six Months Ended June 30,	
(in millions)	2016	2015	2016	2015
Interest income	$1	$—	$4	$1
Foreign currency losses	1	(5)	(4)	(14)
Net gains (losses) on investments	(6)	—	(9)	(1)
Other income (expense), net	—	(3)	(1)	(8)
	$(4)	$(8)	$(10)	$(22)

During the second quarter and first half of 2016, we recognized net losses of $3 million due to investment impairments of certain of our strategic investments. Refer to Note B – Acquisitions and Strategic Investments to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for information regarding our strategic investments.

Tax Rate

Our effective tax rates from continuing operations for the three months ended June 30, 2016 and June 30, 2015, were 47.8% and 2.9%, respectively. For the first half of 2016 and 2015 our effective tax rates from continuing operations were 97.0% and (86.9)%, respectively. The change in our reported tax rate for the second quarter and first half of 2016, as compared to the same periods in 2015, relates primarily to the impact of certain receipts and charges that are taxed at different rates than our effective tax rate, including acquisition-related, contingent consideration, litigation-related and restructuring-related items, as well as the impact of certain discrete tax items.

We are contesting significant proposed adjustments from the Internal Revenue Service (IRS) related to its audit of our transfer pricing methodologies for the 2001 through 2007 tax years. The IRS also proposed similar adjustments for the 2008 through 2010 tax years. We disagree with the methodologies being applied by the IRS and we were scheduled to go to trial in the U.S. Tax Court in late July 2016. On July 19, 2016, we entered a Stipulation of Settled Issues intended to resolve all of the transfer pricing issues, as well as the issues related to our transaction with Abbott. The Stipulation of Settled Issues is contingent upon the IRS Office of Appeals applying the same basis of settlement to all transfer pricing issues for the Company’s 2008, 2009, and 2010 tax years, and if applicable, review by the United States Congress Joint Committee on Taxation. If finalized, payments related to the resolution are expected in the next

12 to 24 months. We believe that our income tax reserves associated with these matters are adequate as of June 30, 2016 and we do not expect to recognize any additional charges related to resolution of this controversy. However, the final resolution of these issues is contingent and if and the Stipulation of Settled Issues is not finalized, it could have a material impact on our financial condition, results of operations, or cash flows.

Refer to Note H – Income Taxes to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for information regarding our tax litigation.

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Critical Accounting Policies and Estimates

Our financial results are affected by the selection and application of accounting policies and methods. In the three months ended June 30, 2016, there were no material changes to the application of critical accounting policies and estimates as described in our most recent Annual Report on Form 10-K.

Goodwill Valuation

We allocate any excess purchase price over the fair value of the net tangible and identifiable intangible assets acquired in a business combination to goodwill. We test our goodwill balances during the second quarter of each year for impairment, or more frequently if indicators are present or changes in circumstances suggest that impairment may exist. We assess goodwill for impairment at the reporting unit level, which is defined as an operating segment or one level below an operating segment, referred to as a component. In 2016 and 2015, we identified six operating segments including Interventional Cardiology, Peripheral Interventions, Rhythm Management, Endoscopy, Urology and Pelvic Health, and Neuromodulation. For purposes of identifying our reporting units, we then assessed whether any components of these segments constitute a business for which discrete financial information is available and where segment management regularly reviews the operating results of that component. We identified Rhythm Management as having two components: Cardiac Rhythm Management and Electrophysiology.

For our 2016 and 2015 annual impairment assessment we identified seven reporting units, which align to our seven core businesses: Interventional Cardiology, Peripheral Interventions, Cardiac Rhythm Management, Electrophysiology, Endoscopy, Urology and Pelvic Health and Neuromodulation. For our 2016 annual impairment assessment we aggregated the Cardiac Rhythm Management and Electrophysiology reporting units, components of the Rhythm Management operating segment, based on the criteria prescribed in FASB ASC Topic 350 - Intangibles - Goodwill and Other. These reporting units were aggregated due to a reorganization that commenced in 2015 that resulted in integrated leadership, shared resources and consolidation of certain sites in 2016.

In performing the goodwill impairment assessment, we utilize both the optional qualitative assessment and the two-step approach prescribed under FASB ASC Topic 350, Intangibles - Goodwill and Other. Beginning in 2016, the qualitative assessment was used for testing certain reporting units where fair value has historically exceeded carrying value by greater than 100%. All other reporting units were tested using the two-step approach described below. The qualitative assessment requires an evaluation of whether it is more likely than not that the fair value of the reporting unit is less than its carrying amount based on an assessment of relevant events including macroeconomic factors, industry and market conditions, cost factors, overall financial performance and other entity-specific factors. After assessing the totality of events, if it is determined that it is not more likely than not that the fair value of the reporting unit is less than its carrying value, the first and second steps of the goodwill impairment test are unnecessary. If it is determined that impairment is more likely than not, then we perform the first step of the two-step impairment test. In 2016, for all reporting units tested using the optional qualitative assessment, we concluded that it was not necessary to perform the first step of the two-step goodwill impairment test. The first step requires a comparison of the carrying value of the reporting units to the fair value of these units.

For our 2016 and 2015 annual impairment assessment, for those reporting units for which a quantitative test was performed, we used only the income approach, specifically the discounted cash flow (DCF) method, to derive the fair value of each of our reporting units in preparing our goodwill impairment assessments. This approach calculates fair value by estimating the after-tax cash flows attributable to a reporting unit and then discounting these after-tax cash flows to a present value using a risk-adjusted discount rate. We selected this method as being the most meaningful in preparing our goodwill assessments because we believe the income approach most appropriately measures our income producing assets. We have considered using the market approach and cost approach but concluded they are not appropriate in valuing our reporting units given the lack of relevant market comparisons available for application of the market approach and the inability to replicate the value of the specific technology-based assets within our

reporting units for application of the cost approach. Therefore, we believe that the income approach represents the most appropriate valuation technique for which sufficient data are available to determine the fair value of our reporting units.

In applying the income approach to our accounting for goodwill, we make assumptions about the amount and timing of future expected cash flows, terminal value growth rates and appropriate discount rates. The amount and timing of future cash flows within our DCF analysis is based on our most recent operational budgets, long range strategic plans and other estimates. The terminal value growth rate is used to calculate the value of cash flows beyond the last projected period in our DCF analysis and reflects our best estimates for stable, perpetual growth of our reporting units. We use estimates of market-participant risk-adjusted weighted-average cost of capital (WACC) as a basis for determining the discount rates to apply to our reporting units' future expected cash flows.

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If the carrying value of a reporting unit exceeds its fair value, we then perform the second step of the goodwill impairment test to measure the amount of impairment loss, if any. If the carrying value of a reporting unit is zero or negative, we evaluate whether it is more likely than not that a goodwill impairment exists. If we determine adverse qualitative factors exist that would indicate it is more likely than not an impairment exists, we then perform the second step of the goodwill test. The second step of the goodwill impairment test compares the estimated fair value of a reporting unit's goodwill to its carrying value.

Although we use consistent methodologies in developing the assumptions and estimates underlying the fair value calculations used in our impairment tests, these estimates are uncertain by nature and can vary from actual results. The use of alternative valuation assumptions, including estimated revenue projections, growth rates, cash flows and discount rates could result in different fair value estimates.

In the second quarter of 2016, we performed our annual goodwill impairment test for all of our reporting units and concluded the fair value of each reporting unit exceeded its carrying value.Because our global Electrophysiology reporting unit was identified as being at higher risk of potential goodwill impairment during our 2015 annual test, it was tested for impairment on a stand-alone basis in the second quarter of 2016, immediately prior to aggregating it with our global Cardiac Rhythm Management reporting unit. The fair value exceeded the carrying value by approximately 36 percent. In comparison, the global Electrophysiology reporting unit had excess fair value of approximately 28 percent as of our 2015 annual test. As of the date of our 2016 annual goodwill impairment test, the aggregated global Electrophysiology and Cardiac Rhythm Management reporting unit (Rhythm Management) had excess fair value over carrying value of approximately 70 percent and held $292 million of allocated goodwill. As such, it was not deemed at higher risk of future impairment. Changes in our reporting units or in the structure of our business as a result of future reorganizations, acquisitions or divestitures of assets or businesses could result in future impairments of goodwill within our reporting units.

Refer to Note C - Goodwill and Other Intangible Assets to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for additional details related to our annual goodwill impairment test performed in the second quarter of 2016.

Liquidity and Capital Resources

Based on our current business plan, we believe our existing balance of cash and cash equivalents, future cash generated from operations and access to capital markets and our revolving credit facility will be sufficient to fund our operations, invest in our infrastructure, pay our legal-related liabilities, fund possible mergers and/or acquisitions and service our existing debt for the next twelve months. On July 12, 2016, we were denied our petition for certiorari by the Maryland Court of Appeals related to our complaint against Mirowski Family Ventures LLC. As a result of the denied petition, we paid $366 million in satisfaction of judgment and interest on July 26, 2016. The payment was funded through cash on hand, cash from our continuing operations and our revolving credit facility. See Note I - Commitments and Contingencies for additional information on the settlement agreement.

As of June 30, 2016, we had $438 million of cash and cash equivalents on hand, comprised of $77 million invested in money market and government funds, $60 million invested in short-term time deposits, and $301 million in interest bearing and non-interest bearing bank accounts. We invest excess cash on hand in short-term financial instruments that earn market interest rates while mitigating principal risk through instrument and counterparty diversification, as well as what we believe to be prudent instrument selection. We limit our direct exposure to securities in any one industry or issuer. We also have full access to our $2.000 billion revolving credit facility and our $300 million credit and security facility secured by our U.S. trade receivables, both described below.

The following provides a summary and description of our net cash inflows (outflows) for the six months ended June 30, 2016 and 2015:

	Six Months Ended June 30,	
(in millions)	2016	2015
Cash provided by (used for) operating activities	$537	$(137)
Cash provided by (used for) investing activities	(150)	(297)
Cash provided by (used for) financing activities	(268)	752

Operating Activities

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During the first half of 2016, cash provided by operating activities was $537 million, as compared to cash used for operating activities of $137 million during the first half of 2015, an increase of $674 million. This increase was primarily driven by a payment of $300 million made to Johnson & Johnson in the first quarter of 2015 as a result of the aggregate $600 million settlement agreement signed on February 13, 2015 to settle the breach of merger agreement lawsuit brought by Johnson & Johnson against Guidant, stemming from our acquisition of Guidant in 2006. As a result of the settlement agreement, Johnson & Johnson dismissed permanently its action without acknowledgment of liability by Guidant. In exchange, we paid $600 million to Johnson & Johnson during the first half of 2015.

Investing Activities

During the first half of 2016, cash used for investing activities primarily included purchases of property, plant and equipment of $138 million and purchases of privately-held equity securities of $36 million, partially offset by proceeds from the sale of one of two buildings located in Quincy, Massachusetts of $29 million. During the first half of 2015, cash used for investing activities primarily included purchases of privately-held equity securities of $140 million, purchases of property, plant and equipment of $92 million, and payments for the acquisitions of businesses, net of cash acquired of $63 million.

Financing Activities

Our cash flows from financing activities in the first half of 2016 and first half of 2015 reflect repayments of debt, payments of acquisition-related contingent consideration, proceeds from, and cash used to net share settle and stock issuances related to our equity incentive programs.

Debt

We had total debt of $5.427 billion as of June 30, 2016 and $5.677 billion as of December 31, 2015. The debt maturity schedule for the significant components of our debt obligations as of June 30, 2016 is as follows:

(in millions)	2016	2017	2018	2019	2020	Thereafter	Total
Senior Notes	$ —	$250	$600	$—	$1,450	$ 2,350	$4,650
Term Loans	—	—	225	150	375	—	750
	$ —	$250	$825	$150	$1,825	$ 2,350	$5,400

Note: The table above does not include unamortized discounts associated with our senior notes, or amounts related to interest rate contracts used to hedge the fair value of certain of our senior notes.

Revolving Credit Facility

On April 10, 2015, we entered into a new $2.000 billion revolving credit facility (the 2015 Facility) with a global syndicate of commercial banks and terminated our previous $2.000 billion revolving credit facility. The 2015 Facility matures on April 10, 2020. Eurodollar and multicurrency loans under the 2015 Facility bear interest at LIBOR plus an interest margin of between 0.900 percent and 1.500 percent, based on our corporate credit ratings and consolidated leverage ratio (1.300 percent as of June 30, 2016). In addition, we are required to pay a facility fee based on our credit ratings, consolidated leverage ratio and the total amount of revolving credit commitment, regardless of usage, under the credit agreement (0.200 percent per year as of June 30, 2016). The 2015 Facility contains covenants which, among other things, require that we maintain a minimum interest coverage ratio of 3.0 times consolidated EBITDA and a maximum leverage ratio of 4.5 times consolidated EBITDA for the first four fiscal quarter-ends following the closing of the AMS Portfolio Acquisition on August 3, 2015, and decreasing to 4.25 times, 4.0 times, and 3.75 times consolidated EBITDA for the next three fiscal quarter-ends after such four fiscal quarter-ends, respectively, and then to 3.50 times for each fiscal quarter-end thereafter. There were no amounts borrowed under our current and prior

revolving credit facilities as of June 30, 2016 or December 31, 2015.

	Covenant Requirement as of June 30, 2016	Actual as of June 30, 2016
Maximum leverage ratio (1)	4.5 times	2.6 times
Minimum interest coverage ratio (2)	3.0 times	8.8 times

(1) Ratio of total debt to consolidated EBITDA, as defined by the credit agreement, for the preceding four consecutive fiscal quarters.

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(2) Ratio of consolidated EBITDA, as defined by the credit agreement, to interest expense for the preceding four consecutive fiscal quarters.

The credit agreement for the 2015 Facility provides for an exclusion from the calculation of consolidated EBITDA, as defined by the credit agreement, through the credit agreement maturity, of any non-cash charges and up to $620 million in restructuring charges and restructuring-related expenses related to our current or future restructuring plans. As of June 30, 2016, we had $523 million of the restructuring charge exclusion remaining. In addition, any cash litigation payments (net of any cash litigation receipts), as defined by the agreement, are excluded from the calculation of consolidated EBITDA and any new debt issued to fund any tax deficiency payments is excluded from consolidated total debt, as defined in the agreement, provided that the sum of any excluded net cash litigation payments and any new debt issued to fund any tax deficiency payments not exceed $2.000 billion in the aggregate. As of June 30, 2016, we had $1.543 billion of the combined legal and debt exclusion remaining.

As of and through June 30, 2016, we were in compliance with the required covenants.

Term Loans

As of June 30, 2016, we had an aggregate of $750 million outstanding under our unsecured term loan facilities and $1.000 billion outstanding as of December 31, 2015. These facilities include an unsecured term loan facility entered into in August 2013 (2013 Term Loan) which had $150 million outstanding as of June 30, 2016 and $250 million outstanding as of December 31, 2015, along with an unsecured term loan credit facility entered into in April 2015 (2015 Term Loan) which had $600 million outstanding as of June 30, 2016 and $750 million outstanding as of December 31, 2015.

Borrowings under the 2013 Term Loan bear interest at LIBOR plus an interest margin between 1.00 percent and 1.75 percent (currently 1.50 percent) based on our corporate credit ratings and consolidated leverage ratio. We repaid $150 million of our 2013 Term Loan facility in the fourth quarter of 2015 and repaid an additional $100 million during the second quarter of 2016. As a result and in accordance with the credit agreement, the outstanding balance of $150 million is the remaining principal amount due at the final maturity date in August 2018. The 2013 Term Loan borrowings are repayable at any time without premium or penalty. Our term loan facility requires that we comply with certain covenants, including financial covenants with respect to maximum leverage and minimum interest coverage, consistent with the 2015 Term Loan facility. The maximum leverage ratio requirement is 4.5 times and our actual leverage ratio as of June 30, 2016 is 2.6 times. The minimum interest coverage ratio requirement is 3.0 times and our actual interest coverage ratio as of June 30, 2016 is 8.8 times.

Our 2015 Term Loan for $750 million was funded on August 3, 2015 and was used to partially fund the AMS Portfolio Acquisition, including the payment of fees and expenses. Term loan borrowings under this facility bear interest at LIBOR plus an interest margin of between 1.00 percent and 1.75 percent (currently 1.50 percent), based on our corporate credit ratings and consolidated leverage ratio. We repaid $150 million of our 2015 Term Loan during the second quarter of 2016. The remaining 2015 Term Loan requires quarterly principal payments of $38 million commencing in the third quarter of 2018, and the remaining principal amount is due at the final maturity date of August 3, 2020. The 2015 Term Loan agreement requires that we comply with certain covenants, including financial covenants with respect to maximum leverage and minimum interest coverage, consistent with our revolving credit facility. The maximum leverage ratio requirement is 4.5 times and our actual leverage ratio as of June 30, 2016 is 2.6 times. The minimum interest coverage ratio requirement is 3.0 times and our actual interest coverage ratio as of June 30, 2016 is 8.8 times.

Senior Notes

We had senior notes outstanding of $4.650 billion as of June 30, 2016 and December 31, 2015. In May 2015, we completed the offering of $1.850 billion in aggregate principal amount of senior notes consisting of $600 million in aggregate principal amount of 2.850% notes due 2020, $500 million in aggregate principal amount of 3.375% notes due 2022 and $750 million in aggregate principal amount of 3.850% notes due 2025. The net proceeds from the offering of the notes, after deducting underwriting discounts and estimated offering expenses, were approximately $1.831 billion. We used a portion of the net proceeds from the senior notes offering to redeem $400 million aggregate principal amount of our 5.500% notes due November 2015 and $600 million aggregate principal amount of our 6.400% notes due June 2016. The remaining senior notes offering proceeds, together with the 2015 Term Loan, were used to fund the AMS Portfolio Acquisition. We recorded a charge of $45 million in interest expense, during the second quarter of 2015, for premiums, accelerated amortization of debt issuance costs, and investor discount costs net of interest rate hedge gains related to the early debt extinguishment.

Our senior notes were issued in public offerings, are redeemable prior to maturity and are not subject to any sinking fund requirements. Our senior notes are unsecured, unsubordinated obligations and rank on parity with each other. These notes are

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effectively junior to borrowings under our credit and security facility, to the extent if borrowed by our subsidiaries and to liabilities of our subsidiaries (see Other Arrangements below).

Other Arrangements

We maintain a $300 million credit and security facility secured by our U.S. trade receivables maturing on June 9, 2017. The credit and security facility requires that we maintain a maximum leverage covenant consistent with our revolving credit facility. The maximum leverage ratio requirement is 4.5 times and our actual leverage ratio as of June 30, 2016 is 2.6 times. We had no borrowings outstanding under this facility as of June 30, 2016 and December 31, 2015.

We have accounts receivable factoring programs in certain European countries that we account for as sales under FASB ASC Topic 860, Transfers and Servicing. These agreements provide for the sale of accounts receivable to third parties, without recourse, of up to approximately $400 million as of June 30, 2016. We have no retained interests in the transferred receivables, other than collection and administrative responsibilities and, once sold, the accounts receivable are no longer available to satisfy creditors in the event of bankruptcy. We de-recognized $169 million of receivables as of June 30, 2016 at an average interest rate of 1.8 percent, and $151 million as of December 31, 2015 at an average interest rate of 2.4 percent.

In addition, we have uncommitted credit facilities with a commercial Japanese bank that provide for borrowings, promissory notes discounting and receivables factoring of up to 21.000 billion Japanese yen (approximately $204 million as of June 30, 2016). We de-recognized $170 million of notes receivable and factored receivables as of June 30, 2016 at an average interest rate of 1.6 percent and $132 million of notes receivable as of December 31, 2015 at an average interest rate of 1.6 percent. De-recognized accounts and notes receivable are excluded from trade accounts receivable, net in the accompanying unaudited condensed consolidated balance sheets.

As of June 30, 2016 we had outstanding letters of credit of $43 million, as compared to $44 million as of December 31, 2015, which consisted primarily of bank guarantees and collateral for workers' compensation insurance arrangements. As of June 30, 2016 and December 31, 2015, none of the beneficiaries had drawn upon the letters of credit or guarantees; accordingly, we did not recognize a related liability for our outstanding letters of credit in our consolidated balance sheets as of June 30, 2016 or December 31, 2015. We believe we will generate sufficient cash from operations to fund these arrangements and intend to fund these arrangements without drawing on the letters of credit.

Equity

During the first half of 2016 and 2015, we received $73 million and $70 million, respectively, in proceeds from stock issuances related to our stock option and employee stock purchase plans. Proceeds from the exercise of employee stock options and employee stock purchases vary from period to period based upon, among other factors, fluctuations in the trading price of our common stock and in the exercise and stock purchase patterns of our employees.

We did not repurchase any shares of our common stock during the six months ended June 30, 2016 and June 30, 2015. As of June 30, 2016, the remaining authorization to repurchase shares under our 2013 share repurchase program was $535 million.

Stock-based compensation expense related to our stock ownership plans was approximately $58 million for the first half of 2016 and $53 million for the first half of 2015.

Contractual Obligations and Commitments

Certain of our acquisitions involve the payment of contingent consideration. See Note B - Acquisitions and Strategic Investments to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for further details regarding the estimated potential amount of future contingent consideration we could be required to pay associated with our acquisitions. There have been no other material changes to our contractual

obligations and commitments as reported in our most recent Annual Report filed on Form 10-K.

Legal Matters

For a discussion of our material legal proceedings see Note I - Commitments and Contingencies to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q and Note I – Commitments and Contingencies to our audited financial statements contained in Item 8 of our most recent Annual Report on Form 10-K.

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Recent Accounting Pronouncements

Information regarding new accounting pronouncements is included in Note M - New Accounting Pronouncements to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q.

Additional Information

Use of Non-GAAP Financial Measures

To supplement our unaudited condensed consolidated financial statements presented on a GAAP basis, we disclose certain non-GAAP financial measures, including adjusted net income (earnings) and adjusted net income (earnings) per share that exclude certain amounts, and adjusted net sales and growth rates that exclude the impact of changes in foreign currency exchange rates. These non-GAAP financial measures are not in accordance with generally accepted accounting principles in the United States.

The GAAP financial measure most directly comparable to adjusted net income is GAAP net income and the GAAP financial measure most directly comparable to adjusted net income per share is GAAP net income per share. To calculate adjusted net sales excluding the impact of changes in foreign currency exchange rates, we convert actual net sales from local currency to U.S. dollars using constant foreign currency exchange rates in the current and prior period. The GAAP financial measure most directly comparable to this non-GAAP financial measure is net sales on a GAAP basis. Reconciliations of each of these non-GAAP financial measures to the corresponding GAAP financial measure are included elsewhere in this Quarterly Report on Form 10-Q.

Management uses these supplemental non-GAAP financial measures to evaluate performance period over period, to analyze the underlying trends in our business, to assess our performance relative to our competitors, and to establish operational goals and forecasts that are used in allocating resources. In addition, management uses these non-GAAP financial measures to further its understanding of the performance of our operating segments. The adjustments excluded from our non-GAAP financial measures are consistent with those excluded from our operating segments' measures of net sales and profit or loss. These adjustments are excluded from the segment measures that are reported to our chief operating decision maker that are used to make operating decisions and assess performance.

We believe that presenting adjusted net income and adjusted net income per share that exclude certain amounts, and adjusted net sales that exclude the impact of changes in foreign currency exchange rates, in addition to the corresponding GAAP financial measures, provides investors greater transparency to the information used by management for its financial and operational decision-making and allows investors to see our results "through the eyes" of management. We further believe that providing this information assists our investors in understanding our operating performance and the methodology used by management to evaluate and measure such performance.

Adjusted net income and adjusted net income per share that exclude certain amounts, and adjusted net sales that exclude the impact of changes in foreign currency exchange rates, are not in accordance with U.S. GAAP and should not be considered in isolation from or as a replacement for the most directly comparable GAAP financial measures. Further, other companies may calculate these non-GAAP financial measures differently than we do, which may limit the usefulness of those measures for comparative purposes.

The following is an explanation of each of the adjustments that management excluded as part of these non-GAAP financial measures for the six months ended June 30, 2016 and 2015, as well as reasons for excluding each of these individual items:

Adjusted Net Income and Adjusted Net Income per Share

Intangible asset impairment charges - This amount represents write-downs of certain intangible asset balances in the first half of 2015. We review intangible assets subject to amortization quarterly to determine if any adverse conditions exist or a change in circumstances has occurred that would indicate impairment and test our indefinite-lived intangible assets at least annually for impairment. If we determine the carrying value of the amortizable intangible asset is not recoverable or we conclude that it is more likely than not that the indefinite-live asset is impaired, we will write the carrying value down to fair value in the period identified. We exclude the impact of impairment charges from management's assessment of operating performance and from our operating segments' measures of profit and loss used for making operating decisions and assessing performance. Accordingly, management has excluded intangible asset impairment charges for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.

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Acquisition- and divestiture-related net charges (credits) - These adjustments may consist of (a) contingent consideration fair value adjustments; (b) gains on previously held investments; (c) purchased and/or funded in-process research and development expenses incurred outside of a business combination; and (d) due diligence, other fees, inventory step up amortization, and integration and exit costs. The contingent consideration adjustments represent accounting adjustments to state contingent consideration liabilities at their estimated fair value. These adjustments can be highly variable depending on the assessed likelihood and amount of future contingent consideration payments. Due diligence, other fees, inventory step-up amortization, and integration and exit costs include legal, tax, severance and other expenses associated with prior and potential future acquisitions that can be highly variable and not representative of ongoing operations. Accordingly, management excluded these amounts for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.

- Restructuring and restructuring-related net charges (credits) - These adjustments represent severance and other direct costs associated with our restructuring programs. These costs are excluded by management in assessing our operating performance, as well as from our operating segments' measures of profit and loss used for making operating decisions and assessing performance. Accordingly, management excluded these costs for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.
- Litigation-related net charges (credits) - These adjustments include certain significant product liability and other litigation-related charges and credits. These amounts are excluded by management in assessing our operating performance, as well as from our operating segments' measures of profit and loss used for making operating decisions and assessing performance. Accordingly, management excluded these amounts for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.
- Debt extinguishment charges - This item represents premiums, accelerated amortization of debt issuance costs and investor discount costs net of interest rate hedge gains related to the early extinguishment of $1.0 billion of public senior notes during the second quarter of 2015. These adjustments are not expected to recur and do not reflect expected ongoing operating results. Accordingly, management excluded these amounts for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.
- Pension termination charges - This item represents charges associated with the termination of the Guidant Retirement Plan, a frozen defined benefit plan. These charges are not expected to recur after 2015 and do not reflect expected ongoing operating results. Accordingly, management has excluded these amounts for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.
- Amortization expense - We record intangible assets at historical cost and amortize them over their estimated useful lives. Amortization expense is excluded from management's assessment of operating performance and is also excluded from our operating segments' measures of profit and loss used for making operating decisions and assessing performance. Accordingly, management has excluded amortization expense for purposes of calculating these non-GAAP financial measures to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.

Adjusted Net Sales Excluding the Impact of Changes in Foreign Currency Exchange Rates

- The impact of changes in foreign currency exchange rates is highly variable and difficult to predict. Accordingly, management excludes the impact of changes in foreign currency exchange rates for purposes of reviewing the net sales and growth rates to facilitate an evaluation of our current operating performance and a comparison to our past operating performance.

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Safe Harbor for Forward-Looking Statements

Certain statements that we may make from time to time, including statements contained in this Quarterly Report on Form 10-Q and information incorporated by reference herein, constitute "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. Forward-looking statements may be identified by words like "anticipate," "expect," "project," "believe," "plan," "may," "estimate," "intend" and similar words. These forward-looking statements are based on our beliefs, assumptions and estimates using information available to us at the time and are not intended to be guarantees of future events or performance. If our underlying assumptions turn out to be incorrect, or if certain risks or uncertainties materialize, actual results could vary materially from the expectations and projections expressed or implied by our forward-looking statements. As a result, readers are cautioned not to place undue reliance on any of our forward-looking statements. Except as required by law, we do not intend to update any forward-looking statements even if new information becomes available or other events occur in the future.

The forward-looking statements in this Quarterly Report on Form 10-Q are based on certain risks and uncertainties, including the risk factors described in "Item 1A. Risk Factors" of this Quarterly Report on Form 10-Q, "Part I, Item 1A. Risk Factors" in our 2015 Annual Report on Form 10-K and the specific risk factors discussed below and in connection with forward-looking statements throughout this Quarterly Report on Form 10-Q, which could cause actual results to vary materially from the expectations and projections expressed or implied by our forward-looking statements. These factors, in some cases, have affected and in the future could affect our ability to implement our business strategy and may cause actual results to differ materially from those contemplated by the forward-looking statements. These additional factors include, among other things, future political, economic, competitive, reimbursement and regulatory conditions; new product introductions; demographic trends; intellectual property; litigation and governmental investigations; financial market conditions; and future business decisions made by us and our competitors, all of which are difficult or impossible to predict accurately and many of which are beyond our control. We caution each reader of this Quarterly Report on Form 10-Q to consider carefully these factors.

The following are some of the important risk factors that could cause our actual results to differ materially from our expectations in any forward-looking statements. For further discussion of these and other risk factors, see "Item 1A. Risk Factors" of this Quarterly Report on Form 10-Q and "Part I, Item 1A. Risk Factors" in our 2015 Annual Report on Form 10-K.

Our Businesses

- Our ability to increase net sales, expand the market and capture market share;

- The volatility of the coronary stent market and our ability to increase our drug-eluting stent systems net sales, including with respect to our SYNERGY™, Promus PREMIER™ and PROMUS® Element™ stent systems, and capture market share;

- The ongoing impact on our business, of physician alignment to hospitals, governmental investigations and audits of hospitals, and other market and economic conditions on the overall number of procedures performed;

- Competitive offerings and related declines in average selling prices for our products, particularly our drug-eluting coronary stent systems and our CRM products;

- The performance of, and physician and patient confidence in, our products and technologies, or those of our competitors;

- The impact and outcome of ongoing and future clinical trials, and market studies undertaken by us, our competitors or other third parties, or perceived product performance of our or our competitors' products;

- Variations in clinical results, reliability or product performance of our and our competitors' products;

- Our ability to acquire or develop, launch and supply new or next-generation products and technologies worldwide and across our businesses in line with our commercialization strategies in a timely and successful manner, including our S-ICD® system and the acquisition and integration of Cosman Medical, Inc., the interventional radiology portfolio of CeloNova Biosciences, the American Medical Systems male urology portfolio and Xlumena, Inc.,

- The effect of consolidation and competition in the markets in which we do business, or plan to do business;

- Disruption in the manufacture or supply of certain components, materials or products, or the failure to timely secure alternative manufacturing or additional or replacement components, materials or products;

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- Our ability to retain and attract key personnel;

- The impact of enhanced requirements to obtain regulatory approval in the U.S. and around the world, including the associated timing and cost of product approval;

- The impact of increased pressure on the availability and rate of third-party reimbursement for our products and procedures in the U.S. and around the world, including with respect to the timing and costs of creating and expanding markets for new products and technologies; and

- Risk associated with counterparty default on our derivative financial instruments.

Regulatory Compliance and Litigation

- The impact of healthcare policy changes and legislative or regulatory efforts in the U.S. and around the world to modify product approval or reimbursement processes, including a trend toward demonstrating clinical outcomes, comparative effectiveness and cost efficiency, as well as the impact of other healthcare reform legislation;

- Risks associated with our regulatory compliance and quality systems and activities in the U.S. and around the world, including meeting regulatory standards applicable to manufacturing and quality processes;

- Our ability to minimize or avoid future field actions or FDA warning letters relating to our products and processes and the ongoing inherent risk of potential physician advisories related to medical devices;

- The impact of increased scrutiny of and heightened global regulatory enforcement facing the medical device industry arising from political and regulatory changes, economic pressures or otherwise, including under U.S. Anti-Kickback Statute, U.S. False Claims Act and similar laws in other jurisdictions; U.S. Foreign Corrupt Practices Act (FCPA) and/or similar laws in other jurisdictions, and U.S. and foreign export control, trade embargo and customs laws;

- Costs and risks associated with litigation;

- The effect of our litigation and risk management practices, including self-insurance, and compliance activities on our loss contingencies, legal provision and cash flows;

- The impact of, diversion of management attention as a result of, and costs to cooperate with, litigate and/or resolve, governmental investigations and our class action, product liability, contract and other legal proceedings; and

- Risks associated with a failure to protect our intellectual property rights and the outcome of patent litigation.

Innovation and Certain Growth Initiatives

- The timing, size and nature of our strategic growth initiatives and market opportunities, including with respect to our internal research and development platforms and externally available research and development platforms and technologies, and the ultimate cost and success of those initiatives and opportunities;

- Our ability to complete planned clinical trials successfully, obtain regulatory approvals and launch new and next generation products in a timely manner consistent with cost estimates, including the successful completion of in-process projects from in-process research and development;

- Our ability to identify and prioritize our internal research and development project portfolio and our external investment portfolio on profitable revenue growth opportunities as well as to keep them in line with the estimated timing and costs of such projects and expected revenue levels for the resulting products and technologies;

- Our ability to successfully develop, manufacture and market new products and technologies in a timely manner and the ability of our competitors and other third parties to develop products or technologies that render our products or technologies noncompetitive or obsolete;

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- The impact of our failure to succeed at or our decision to discontinue, write-down or reduce the funding of any of our research and development projects, including in-process projects from in-process research and development, in our growth adjacencies or otherwise;

- Dependence on acquisitions, alliances or investments to introduce new products or technologies and to enter new or adjacent growth markets, and our ability to fund them or to fund contingent payments with respect to those acquisitions, alliances and investments; and

- The failure to successfully integrate and realize the expected benefits from the strategic acquisitions, alliances and investments we have consummated or may consummate in the future.

International Markets

- Our dependency on international net sales to achieve growth, including in emerging markets;

- The impact of changes in our international structure and leadership;

- Risks associated with international operations and investments, including the timing and collectibility of customer payments, political and economic conditions, protection of our intellectual property, compliance with established and developing U.S. and foreign legal and regulatory requirements, including FCPA and similar laws in other jurisdictions and U.S. and foreign export control, trade embargo and customs laws, as well as changes in reimbursement practices and policies;

- Our ability to maintain or expand our worldwide market positions in the various markets in which we compete or seek to compete, including through investments in product diversification and emerging markets such as Brazil, Russia, India and China;

- Our ability to execute and realize anticipated benefits from our investments in emerging markets; and

- The potential effect of foreign currency fluctuations and interest rate fluctuations on our net sales, expenses and resulting margins.

Liquidity

- Our ability to generate sufficient cash flow to fund operations, capital expenditures, global expansion initiatives, any litigation settlements and judgments, share repurchases and strategic investments and acquisitions as well as maintaining our investment grade ratings and managing our debt levels and covenant compliance;

- Our ability to access the public and private capital markets when desired and to issue debt or equity securities on terms reasonably acceptable to us;

- The unfavorable resolution of open tax matters, exposure to additional tax liabilities and the impact of changes in U.S. and international tax laws;

- The impact of examinations and assessments by domestic and international taxing authorities on our tax provision, financial condition or results of operations;

- The impact of goodwill and other intangible asset impairment charges, including on our results of operations; and

- Our ability to collect outstanding and future receivables and/or sell receivables under our factoring programs.

Cost Reduction and Optimization Initiatives

- Risks associated with significant changes made or expected to be made to our organizational and operational structure, pursuant to our 2016 Restructuring plan and the completion of our 2014 Restructuring Plan, as well as any further restructuring or optimization plans we may undertake in the future, and our ability to recognize benefits and cost reductions from such programs; and

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- Business disruption and employee distraction as we execute our global compliance program, restructuring and optimization plans and divestitures of assets or businesses and implement our other strategic and cost reduction initiatives.

Rule 10b5-1 Trading Plans by Executive Officers

Periodically, certain of our executive officers adopt written stock trading plans in accordance with Rule 10b5-1 under the Exchange Act and our own Stock Trading Policy (a Rule 10b5-1 Trading Plan). A Rule 10b5-1 Trading Plan is a written document that pre-establishes the amount, prices and dates (or formulas for determining the amounts, prices and dates) of future purchases or sales of our stock, including shares issued upon exercise of stock options or vesting of deferred stock units. These plans are entered into at a time when the person is not in possession of material non-public information about Boston Scientific Corporation.

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ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

We develop, manufacture and sell medical devices globally and our earnings and cash flows are exposed to market risk from changes in currency exchange rates and interest rates. We address these risks through a risk management program that includes the use of derivative financial instruments. We operate the program pursuant to documented corporate risk management policies. We do not enter derivative transactions for speculative purposes. Gains and losses on derivative financial instruments substantially offset losses and gains on underlying hedged exposures. Furthermore, we manage our exposure to counterparty risk on derivative instruments by entering into contracts with a diversified group of major financial institutions and by actively monitoring outstanding positions.

Our currency risk consists primarily of foreign currency denominated firm commitments, forecasted foreign currency denominated intercompany and third-party transactions and net investments in certain subsidiaries. We use both nonderivative (primarily European manufacturing operations) and derivative instruments to manage our earnings and cash flow exposure to changes in currency exchange rates. We had currency derivative instruments outstanding in the contract amount of $4.865 billion as of June 30, 2016 and $3.547 billion as of December 31, 2015. We recorded $144 million of other assets and $199 million of other liabilities to recognize the fair value of these derivative instruments as of June 30, 2016, as compared to $237 million of other assets and $23 million of other liabilities as of December 31, 2015. A ten percent appreciation in the U.S. dollar’s value relative to the hedged currencies would increase the derivative instruments’ fair value by $303 million as of June 30, 2016 and $155 million as of December 31, 2015. A ten percent depreciation in the U.S. dollar’s value relative to the hedged currencies would decrease the derivative instruments’ fair value by $326 million as of June 30, 2016 and by $189 million as of December 31, 2015. Any increase or decrease in the fair value of our currency exchange rate sensitive derivative instruments would be substantially offset by a corresponding decrease or increase in the fair value of the hedged underlying asset, liability or forecasted transaction, resulting in minimal impact on our consolidated statements of operations.

Our interest rate risk relates primarily to U.S. dollar borrowings partially offset by U.S. dollar cash investments. We have historically used interest rate derivative instruments to manage our earnings and cash flow exposure to changes in interest rates. We had no interest rate derivative instruments outstanding as of June 30, 2016. As of June 30, 2016, $4.673 billion of our outstanding debt obligations were at fixed interest rates, representing approximately 86 percent of our total debt.

Refer to Note D – Fair Value Measurements to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q for further information regarding our derivative financial instruments.

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ITEM 4. CONTROLS AND PROCEDURES

Evaluation of Disclosure Controls and Procedures

Our management, with the participation of our Chief Executive Officer (CEO), and our Chief Financial Officer (CFO), evaluated the effectiveness of our disclosure controls and procedures as of June 30, 2016 pursuant to Rule 13a-15(b) of the Securities Exchange Act of 1934, as amended (the Exchange Act). Disclosure controls and procedures are designed to ensure that material information required to be disclosed by us in the reports that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms and that such material information is accumulated and communicated to our management, including our CEO and CFO, as appropriate, to allow timely decisions regarding required disclosure. Based on their evaluation, our CEO and CFO concluded that, as of June 30, 2016, our disclosure controls and procedures were effective.

Changes in Internal Control Over Financial Reporting

During the six month period ended June 30, 2016, there were no changes in our internal control over financial reporting that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

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PART II
OTHER INFORMATION

ITEM 1. LEGAL PROCEEDINGS
See Note H – Income Taxes and Note I – Commitments and Contingencies to our unaudited condensed consolidated financial statements contained in Item 1 of this Quarterly Report on Form 10-Q, which is incorporated herein by reference.

ITEM 1A. RISK FACTORS

In addition to the information set forth below and other information contained elsewhere in this report, you should carefully consider the factors discussed in "Part I, Item 1A. Risk Factors" in our most recent Annual Report filed on Form 10-K, which could materially affect our business, financial condition or future results.

We are subject to a number of market, business, financial, legal and regulatory risks and uncertainties with respect to our international operations that could have a material impact on our business, financial condition or results of operations.

International net sales accounted for approximately 43 percent of our global net sales in 2015, with sales from emerging markets accounting for approximately 10 percent. An important part of our growth strategy is to continue pursuing growth opportunities in net sales and market share outside of the U.S. by expanding global presence, including in emerging markets. Our international operations are subject to a number of market, business and financial risks and uncertainties, including those related to political and economic instability; foreign currency exchange and interest rate fluctuations; competitive product offerings; local changes in health care financing and payment systems and health care delivery systems; local product preferences and requirements, including preferences for local manufacturers; workforce instability; less intellectual property protection in certain countries than exists in the United States; and, in certain foreign countries, longer accounts receivable cycles. Such risks and uncertainties may adversely impact our ability to implement our growth strategy in these markets and, as a result, our sales growth, market share and operating profits from our international operations may be adversely affected.

Our international operations are subject to established and developing legal and regulatory requirements for medical devices in each country in which our products are marketed and sold. Most foreign countries have medical device regulations. Further, most countries outside of the U.S. require product approvals be renewed or recertified on a regular basis in order to continue to be marketed and sold there. In addition, several countries that previously did not have regulatory requirements for medical devices have established such requirements in recent years and other countries have expanded, or plan to expand, on existing regulations, including requiring local clinical data in addition to global clinical data. These factors have caused or may cause us to experience more uncertainty, risk, expense and delay in commercializing products in certain foreign jurisdictions, which could affect our ability to obtain approvals for our products in those jurisdictions and adversely impact our net sales, market share and operating profits from our international operations.

Further, international markets are affected by economic pressure to contain healthcare costs, which can lead to lower reimbursement rates for either our products directly or procedures in which are our products are used. Governments and payers may also institute changes in health care delivery systems that may reduce funding for services or encourage greater scrutiny of health care costs. In addition, certain international markets may also be affected by foreign government efforts to reference reimbursement rates in other countries. All of these types of changes may ultimately reduce selling prices of our products or reduce the number of procedures in which our products are used, which may adversely impact our net sales, market share and operating profits from our international operations.

In addition, our international operations are subject to other established and developing U.S. and foreign legal and regulatory requirements, including the U.S. Foreign Corrupt Practices Act (FCPA) and/or similar laws in other countries; and U.S. and foreign import and export controls and licensing requirements, trade protection and embargo measures and customs laws. Global businesses, including those in the medical device industry, are facing increasing scrutiny of, and heightened enforcement efforts with respect to, their international operations. Any alleged or actual failure to comply with legal and regulatory requirements may subject us to government scrutiny, civil and/or criminal proceedings, sanctions and other liabilities, which may have a material adverse effect on our international operations, financial condition, results of operations and/or liquidity.

Following a referendum in June 2016 in which voters in the United Kingdom (UK) approved an exit from the EU, the UK government is expected to initiate a process to withdraw from the EU (often referred to as “Brexit”) and begin negotiating the terms of the UK’s future relationship with the EU. A withdrawal could, among other outcomes, result in the deterioration of

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economic conditions, volatility in currency exchange rates, and increased regulatory complexities. These outcomes may adversely affect our business, financial condition or results of operations.

Any significant changes in the political and economic, financial, competitive, legal and regulatory or reimbursement conditions where we conduct, or plan to expand, our international operations may have a material impact on our business, financial condition or results of operations.

ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

None.

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ITEM 6. EXHIBITS (* documents filed or furnished with this report, # compensatory plans or arrangements)

31.1*	Certification of the Chief Executive Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
31.2*	Certification of the Chief Financial Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
32.1*	Certification Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, Chief Executive Officer
32.2*	Certification Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, Executive Vice President and Chief Financial Officer
101*	Interactive Data Files Pursuant to Rule 405 of Regulation S-T: (i) the Condensed Consolidated Statements of Operations for the three and six months ended June 30, 2016 and 2015, (ii) the Condensed Consolidated Statements of Comprehensive Income for the three and six months ended June 30, 2016 and 2015, (iii) the Condensed Consolidated Balance Sheets as of June 30, 2016 and December 31, 2015, (iv) the Condensed Consolidated Statements of Cash Flows for the six months ended June 30, 2016 and 2015 and (v) the notes to the Condensed Consolidated Financial Statements.

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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 on August 3, 2016.

BOSTON SCIENTIFIC CORPORATION

By: /s/ Daniel J. Brennan

Name: Daniel J. Brennan

Title: Executive Vice President and Chief Financial Officer