UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 10-Q

(Mark One)

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

For the quarterly period ended June 30, 2017

OR

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

to

For the transition period from

Commission File No. 1-2189

ABBOTT LABORATORIES

An Illinois Corporation

I.R.S. Employer Identification No. 36-0698440

100 Abbott Park Road Abbott Park, Illinois 60064-6400

Telephone: (224) 667-6100

Indicate by check mark whether the registrant: (I) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of l934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes x No o

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§ 229.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). Yes x No o

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

Large Accelerated Filer x

Accelerated Filer o

Non-Accelerated Filer o (Do not check if a smaller reporting company)

Smaller reporting company o

Emerging growth company o

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. 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 x

As of June 30, 2017, Abbott Laboratories had 1,737,443,264 common shares without par value outstanding.

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Abbott Laboratories

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Abbott Laboratories and Subsidiaries Condensed Consolidated Statement of Earnings (Unaudited)

(dollars in millions except per share data; shares in thousands)

	Three Months Ended June 30						Six Months Ended June 30					
		2017		2016		2017		2016				
Net sales	\$	6,637	\$	5,333	\$	12,972	\$	10,218				
Cost of products sold, excluding amortization of intangible												
assets		3,173		2,287		6,217		4,427				
Amortization of intangible assets		392		145		914		289				
Research and development		513		348		1,060		727				
Selling, general and administrative		2,132		1,737		4,556		3,435				
Total operating cost and expenses		6,210		4,517		12,747		8,878				
Operating earnings		427		816		225		1,340				
Interest expense		214		103		440		161				
Interest (income)		(31)		(20)		(53)		(53)				
Net foreign exchange loss (gain)		(12)		10		(28)		488				
Other expense (income), net		(39)		8		(1,165)		27				
Earnings from continuing operations before taxes		295		715		1,031		717				
Taxes on earnings from continuing operations		25		116		375		62				
Earnings from continuing operations		270		599		656		655				
Earnings from discontinued operations, net of tax		13		16		46		260				
Gain on sale of discontinued operations, net of tax		_		_		_		16				
Net earnings from discontinued operations, net of tax		13		16		46		276				
Net Earnings	\$	283	\$	615	\$	702	\$	931				
Basic Earnings Per Common Share —												
Continuing operations	\$	0.15	\$	0.40	\$	0.37	\$	0.44				
Discontinued operations	Ψ	0.01	Ψ	0.01	Ψ	0.03	Ψ	0.19				
Net earnings	\$	0.16	\$	0.41	\$	0.40	\$	0.63				
Diluted Earnings Per Common Share —												
Continuing operations	\$	0.15	\$	0.40	\$	0.37	\$	0.44				
Discontinued operations	Ψ	0.13	Ψ	0.01	Ψ	0.03	Ψ	0.19				
Net earnings	\$	0.16	\$	0.41	\$	0.40	\$	0.63				
- 100 Carrings	Ψ	0.10	Ψ	0.41	Ψ	0.40	Ψ	0.03				
Cash Dividends Declared Per Common Share	\$	0.265	\$	0.26	\$	0.53	\$	0.52				

Average Number of Common Shares Outstanding Used for Basic Earnings Per Common Share	1,740,524	1,474,504	1,734,008	1,476,161
Dilutive Common Stock Options	8,359	5,988	8,099	6,165
Average Number of Common Shares Outstanding Plus Dilutive Common Stock Options	1,748,883	1,480,492	1,742,107	1,482,326
Outstanding Common Stock Options Having No Dilutive Effect	5,258	5,673	5,258	5,673

The accompanying notes to the condensed consolidated financial statements are an integral part of this statement.

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Abbott Laboratories and Subsidiaries Condensed Consolidated Statement of Comprehensive Income (Unaudited) (dollars in millions)

	Three Months Ended June 30				Six Montl June			
		2017		2016	2017		2016	
Net Earnings	\$	283	\$	615	\$ 702	\$	931	
Foreign currency translation gain (loss) adjustments		288		(104)	821		317	
Net actuarial gains (losses) and amortization of net actuarial (losses) and prior								
service (cost) and credits, net of taxes of \$11 and \$23 in 2017 and \$(12) and								
\$(3) in 2016		29		(47)	63		(29)	
Unrealized gains (losses) on marketable equity securities, net of taxes of \$7 and								
\$60 in 2017 and nil in 2016		2		(213)	82		(756)	
Net (losses) for derivative instruments designated as cash flow hedges and								
other, net of taxes of \$(15) and \$(39) in 2017 and \$(2) and \$(24) in 2016		(37)		(8)	(102)		(97)	
Other comprehensive income (loss)		282		(372)	864		(565)	
Comprehensive Income	\$	565	\$	243	\$ 1,566	\$	366	
						_		

		June 30, 2017		ecember 31, 2016
Supplemental Accumulated Other Comprehensive Income (Loss) Information, net of tax:	<u> </u>			
Cumulative foreign currency translation (loss) adjustments	\$	(3,996)	\$	(4,959)
Net actuarial (losses) and prior service cost and credits		(2,215)		(2,284)
Cumulative unrealized gains (losses) on marketable equity securities		13		(69)
Cumulative (losses) gains on derivative instruments designated as cash flow hedges and other		(52)		49
Accumulated other comprehensive income (loss)	\$	(6,250)	\$	(7,263)

The accompanying notes to the condensed consolidated financial statements are an integral part of this statement.

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Abbott Laboratories and Subsidiaries Condensed Consolidated Balance Sheet (Unaudited) (dollars in millions)

	June 30, 2017		ecember 31, 2016
Assets			
Current Assets:			
Cash and cash equivalents	\$ 9,675	\$	18,620
Short-term investments	160		155
Trade receivables, less allowances of \$265 in 2017 and \$250 in 2016	4,633		3,248
Inventories:			
Finished products	2,355		1,624
Work in process	450		294
Materials	798		516
Total inventories	3,603		2,434
Prepaid expenses and other receivables	1,912		1,806
Current assets held for disposition	_		513
Total Current Assets	19,983		26,776
Investments	1,545		2,947
Property and equipment, at cost	14,476		12,366
Less: accumulated depreciation and amortization	7,190		6,661
Net property and equipment	7,286		5,705

Intangible assets, net of amortization	18,653	4,539
Goodwill	22,132	7,683
Deferred income taxes and other assets	1,552	2,263
Non-current assets held for disposition	_	2,753
	\$ 71,151	\$ 52,666
Liabilities and Shareholders' Investment		
Current Liabilities:		
Short-term borrowings	\$ 218	\$ 1,322
Trade accounts payable	1,667	1,178
Salaries, wages and commissions	905	752
Other accrued liabilities	3,357	2,581
Dividends payable	461	391
Income taxes payable	192	188
Current portion of long-term debt	3	3
Current liabilities held for disposition	_	245
Total Current Liabilities	 6,803	6,660
Long-term debt	 23,810	20,681
Post-employment obligations, deferred income taxes and other long-term liabilities	8,750	4,549
Non-current liabilities held for disposition	_	59
Commitments and Contingencies		
Shareholders' Investment:		
Preferred shares, one dollar par value Authorized — 1,000,000 shares, none issued	_	_
Common shares, without par value Authorized — 2,400,000,000 shares		
Issued at stated capital amount — Shares: 2017: 1,962,732,172; 2016: 1,707,475,455	23,012	13,027
Common shares held in treasury, at cost — Shares: 2017: 225,288,908; 2016: 234,606,250	(10,362)	(10,791)
Earnings employed in the business	25,202	25,565
Accumulated other comprehensive income (loss)	(6,250)	(7,263)
Total Abbott Shareholders' Investment	 31,602	20,538
Noncontrolling Interests in Subsidiaries	186	179
Total Shareholders' Investment	31,788	20,717
	\$ 71,151	\$ 52,666

The decompanying notes to the condensed consolidated intalicial statements are an integral part of this statement.

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Repayments of long-term debt

Payment of debt issuance costs

Purchases of common shares

Payment of contingent consideration

Proceeds from stock options exercised

Condensed Consolidated Statement of Cash Flows (Unaudited) (dollars in millions)

Abbott Laboratories and Subsidiaries

	Six Months E	nded June	30
	2017		2016
Cash Flow From (Used in) Operating Activities:			
Net earnings	\$ 702	\$	931
Adjustments to reconcile net earnings to net cash from operating activities -			
Depreciation	508		405
Amortization of intangible assets	914		289
Share-based compensation	263		214
Impact of currency devaluation	_		477
Amortization of inventory step-up	822		_
Gain on sale of businesses	(1,151)		(25
Trade receivables	(56)		(150
Inventories	(127)		(149
Other, net	50		(1,176
Net Cash From Operating Activities	1,925		816
Cash Flow From (Used in) Investing Activities:			
Acquisitions of property and equipment	(527)		(490
Acquisitions of businesses and technologies, net of cash acquired	(13,027)		(7
Proceeds from business dispositions	5,471		25
Proceeds from the sale of Mylan N.V. shares	1,924		_
Sales (purchases) of other investment securities, net	(28)		(800
Other	27		28
Net Cash (Used in) Investing Activities	(6,160)		(1,244
Cash Flow From (Used in) Financing Activities:			
Net (repayments of) short-term debt and other	(1,429)		(28:

(2,507)

(13)

(98)

186

(10)

(132) (25) (520)

130

Dividends paid Net Cash (Used in) Financing Activities	(922) (4,783)	(769) (1,611)
Effect of exchange rate changes on cash and cash equivalents	73	(384)
Net Decrease in Cash and Cash Equivalents Cash and Cash Equivalents, Beginning of Year	(8,945) 18,620	(2,423) 5,001
Cash and Cash Equivalents, End of Period	\$ 9,675	\$ 2,578

The accompanying notes to the condensed consolidated financial statements are an integral part of this statement.

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Abbott Laboratories and Subsidiaries

Notes to the Condensed Consolidated Financial Statements

June 30, 2017

(Unaudited)

Note 1 — Basis of Presentation

The accompanying unaudited, condensed consolidated financial statements have been prepared pursuant to rules and regulations of the Securities and Exchange Commission and, therefore, do not include all information and footnote disclosures normally included in audited financial statements. However, in the opinion of management, all adjustments (which include only normal adjustments) necessary to present fairly the results of operations, financial position and cash flows have been made. It is suggested that these statements be read in conjunction with the financial statements included in Abbott's Annual Report on Form 10-K for the year ended December 31, 2016. The consolidated financial statements include the accounts of the parent company and subsidiaries, after elimination of intercompany transactions.

In March 2016, the Financial Accounting Standards Board (FASB) issued Accounting Standards Update (ASU) 2016-09, *Improvements to Employee Share-Based Payment Accounting*. ASU 2016-09 modifies several aspects of the accounting for share-based payment transactions, including the accounting for income taxes and classification on the statement of cash flows. Abbott adopted the standard in the first quarter of 2017 and the following changes were made to the presentation of Abbott's financial statements:

- · All excess tax benefits or tax deficiencies are now recognized as income tax benefit or expense as applicable. Previously, Abbott recorded the benefits to Shareholders' Investment. The tax benefit recorded in Abbott's Condensed Consolidated Statement of Earnings for the second quarter and first six months of 2017 were \$25 million and \$63 million, respectively. The standard does not permit retrospective presentation of this benefit in prior years.
- The tax benefit or deficiency is required to be classified as an operating activity in the statement of cash flows. Previously, it was required to be classified within financing activities. Abbott has adopted this standard on a prospective basis and has not revised the classification of the excess tax benefit in the prior year's Condensed Consolidated Statement of Cash Flows.

Note 2 — Discontinued Operations

On February 27, 2015, Abbott completed the sale of its developed markets branded generics pharmaceuticals business to Mylan Inc. (Mylan) for 110 million shares (or approximately 22%) of a newly formed entity (Mylan N.V.) that combined Mylan's existing business and Abbott's developed markets branded generics pharmaceuticals business. Mylan N.V. is publicly traded. The shareholder agreement with Mylan N.V. includes voting and other restrictions that prevent Abbott from exercising significant influence over the operating and financial policies of Mylan N.V.

In April 2015, Abbott sold 40.25 million of the 110 million ordinary shares of Mylan N.V. As a result of this sale, Abbott's ownership interest in Mylan N.V. decreased to approximately 14%.

In March 2017, Abbott sold 44 million ordinary shares of Mylan N.V. and received \$1.685 billion in proceeds. In June 2017, Abbott sold an additional 6 million ordinary shares of Mylan N.V. and received \$239 million in proceeds. Abbott recorded an immaterial pre-tax gain in the first six months of 2017, which was recognized in the Other expense (income), net line of the Condensed Consolidated Statement of Earnings. As a result of these share sales, Abbott's ownership interest in Mylan N.V. decreased from approximately 14% to approximately 3.7%.

On February 10, 2015, Abbott completed the sale of its animal health business to Zoetis Inc. Abbott received cash proceeds of \$230 million and reported an after-tax gain on the sale of approximately \$130 million in the first quarter of 2015. In the first quarter of 2016, Abbott received an additional \$25 million of proceeds related to the expiration of a holdback agreement associated with the sale of this business and reported an after-tax gain on the sale of discontinued operations of \$16 million.

On January 1, 2013, Abbott completed the separation of AbbVie Inc. (AbbVie), which was formed to hold Abbott's research-based proprietary pharmaceuticals business. Abbott has retained all liabilities for all U.S. federal and foreign income taxes on income prior to the separation, as well as certain non-income related taxes attributable to AbbVie's business prior to the separation. AbbVie generally will be liable for all other taxes attributable to its business. Earnings from discontinued operations, net of tax of \$46 million and \$260 million in the first six months of 2017 and 2016 were driven primarily by the recognition of net tax benefits as a result of the resolution of various tax positions related to AbbVie's operations for years prior to the separation.

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Note 3 — Assets and Liabilities Held for Disposition

In September 2016, Abbott announced that it entered into a definitive agreement to sell Abbott Medical Optics (AMO), its vision care business, to Johnson & Johnson for \$4.325 billion in cash, subject to customary purchase price adjustments for cash, debt and working capital. The decision to sell AMO reflects Abbott's proactive shaping of its portfolio in line with its strategic priorities. In February 2017, Abbott completed the sale of AMO to Johnson & Johnson and recognized a pre-tax gain of \$1.151 billion, which was reported in the Other expense (income), net line of the Condensed Consolidated Statement of Earnings in the first six months of 2017. Abbott recorded an after-tax gain of \$721 million in the first six months of 2017 related to the sale of AMO.

The operating results of AMO up to the date of sale continued to be included in Earnings from Continuing Operations as they did not qualify for reporting as discontinued operations. For the three months ended June 30, 2017 and 2016, the AMO earnings before taxes included in Abbott's consolidated earnings were nil and \$13 million, respectively. For the first six months ended June 30, 2017 and 2016, the AMO losses before taxes included in Abbott's consolidated earnings were \$18 million and \$44 million, respectively. The following assets and liabilities of this business were reported as held for disposition in Abbott's Condensed Consolidated Balance Sheet as of December 31, 2016:

(in millions)	Dec	ember 31, 2016
Trade receivables, net	\$	222
Total inventories		240
Prepaid expenses and other current assets		51
Current assets held for disposition		513
Net property and equipment		247
Intangible assets, net of amortization		529
Goodwill		1,966
Deferred income taxes and other assets		11
Non-current assets held for disposition		2,753
Total assets held for disposition	\$	3,266
Trade accounts payable	\$	71
Salaries, wages, commissions and other accrued liabilities		174
Current liabilities held for disposition		245
Post-employment obligations, deferred income taxes and other long-term liabilities		59
Total liabilities held for disposition	\$	304

Note 4 — Supplemental Financial Information

Shares of unvested restricted stock that contain non-forfeitable rights to dividends are treated as participating securities and are included in the computation of earnings per share under the two-class method. Under the two-class method, net earnings are allocated between common shares and participating securities. Earnings from Continuing Operations allocated to common shares for the three months ended June 30, 2017 and 2016 were \$269 million and \$597 million, respectively and for the six months ended June 30, 2017 and 2016 were \$653 million and \$652 million, respectively. Net earnings allocated to common shares for the three months ended June 30, 2017 and 2016 were \$281 million and \$612 million, respectively, and for the six months ended June 30, 2017 and 2016 were \$698 million and \$927 million, respectively.

The Other, net line in Net cash from operating activities in the Condensed Consolidated Statement of Cash Flows for the first six months of 2017 and 2016 includes the effects of contributions to defined benefit plans of \$321 million and \$524 million, respectively, and to the post-employment medical and dental benefit plans of \$11 million and \$9 million, respectively. The first six months of 2017 also includes the impact of approximately \$430 million of tax expense related to business dispositions, which has not yet been paid, and is taxed at a discrete tax rate. The first six months of 2016 included the non-cash impact of approximately \$410 million of net tax benefits primarily associated with the resolution of various tax positions from prior years, as well as cash taxes paid of approximately \$140 million related to the disposition of businesses. The foreign currency loss related to Venezuela in the first six months of 2016 reduced Abbott's cash by approximately \$410 million and is shown on the Effect of exchange rate changes on cash and cash equivalents line within the Condensed Consolidated Statement of Cash Flows.

Since January 2010, Venezuela has been designated as a highly inflationary economy under U.S. GAAP. In 2014 and 2015, the government of Venezuela operated multiple mechanisms to exchange bolivars into U.S. dollars. These mechanisms included the

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CENCOEX, SICAD, and SIMADI rates, which stood at 6.3, 13.5, and approximately 200, respectively, at December 31, 2015. In 2015, Abbott continued to use the CENCOEX rate of 6.3 Venezuelan bolivars to the U.S. dollar to report the results, financial position, and cash flows related to its operations in Venezuela since Abbott continued to qualify for this exchange rate to pay for the import of various products into Venezuela.

On February 17, 2016, the Venezuelan government announced that the three-tier exchange rate system would be reduced to two rates renamed the DIPRO and DICOM rates. The DIPRO rate is the official rate for food and medicine imports and was adjusted from 6.3 to 10 bolivars per U.S. dollar. The DICOM rate is a floating market rate published daily by the Venezuelan central bank, which at the end of the first quarter of 2016 was approximately 263 bolivars per U.S. dollar. As a result of decreasing government approvals to convert bolivars to U.S. dollars to pay for intercompany accounts, as well as the accelerating deterioration of economic conditions in the country, Abbott concluded that it was appropriate to move to the DICOM rate at the end of the first quarter of 2016. As a result, Abbott recorded a foreign currency exchange loss of \$477 million in the first quarter of 2016 to revalue its net monetary assets in Venezuela. Abbott is continuing to use the DICOM rate to report the results of operations and to remeasure net monetary assets for Venezuela at the end of each quarter. As of June 30, 2017, Abbott's Venezuelan operations represented approximately 0.01% of Abbott's consolidated assets and any additional foreign currency losses related to Venezuela are not expected to be material.

The components of long-term investments as of June 30, 2017 and December 31, 2016 are as follows:

Long-term Investments (in millions)	June 30, 2017			December 31, 2016
Equity securities	\$	1,482	\$	2,906
Other		63		41
Total	\$	1,545	\$	2,947

As discussed in Note 2, in the first six months of 2017, Abbott sold 50 million ordinary shares of Mylan N.V., thereby reducing Abbott's investment in equity securities by approximately \$1.9 billion.

Abbott's equity securities as of June 30, 2017, include approximately \$348 million of investments in mutual funds that are held in a rabbi trust and were acquired as part of the St. Jude Medical, Inc. (St. Jude Medical) business acquisition. These investments, which are specifically designated as available for the purpose of paying benefits under a deferred compensation plan, are not available for general corporate purposes and are subject to creditor claims in the event of insolvency.

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Note 5 — Changes in Accumulated Other Comprehensive Income (Loss)

The changes in accumulated other comprehensive income (loss), net of income taxes, are as follows:

		Three Months Ended June 30														
	Cumulative Foreign Currency Translation Adjustments					Net Actuarial Losses and Prior Service Costs and Credits				Cumulative Unrealized Gains (Losses) on Marketable Equity Securities				Cumulative Gains (Losses) on Derivative Instruments Designated as Cash Flow Hedges		
(in millions)		2017		2016		2017		2016		2017		2016		2017		2016
Balance at March 31	\$	(4,284)	\$	(4,408)	\$	(2,244)	\$	(1,940)	\$	11	\$	(478)	\$	(15)	\$	(25)
Other comprehensive income (loss) before reclassifications		288		(104)		_		(62)		2		(213)		(38)		11
Amounts reclassified from accumulated other comprehensive income		_		_		29		15		_		_		1		(19)
Net current period comprehensive income (loss)		288		(104)		29		(47)		2		(213)		(37)		(8)
. ,																(8)
Balance at June 30	\$	(3,996)	\$	(4,512)	\$	(2,215)	\$	(1,987)	\$	13	\$	(691)	\$	(52)	\$	(33)

								Six Months En	ded .	June 30						
	Cumulative Foreign Currency Translation Adjustments					Net Ac Losses a Service C Cre	nd I Cost	Prior s and		Cumu Unrealize (Losse Marketab Secur	d Ga s) or le Ec	ains 1 Juity	 Cumulative Gains (Losses) on Derivative Instruments Designated as Cash Flow Hedges			
(in millions)	_	2017	_	2016	_	2017	_	2016		2017		2016	 2017	_	2016	
Balance at December 31, 2016 and 2015	\$	(4,959)	\$	(4,829)	\$	(2,284)	\$	(1,958)	\$	(69)	\$	65	\$ 49	\$	64	
Impact of business																
dispositions		142		_		6		_		_		_	1		_	
Other comprehensive income (loss) before																
reclassifications		821		317		_		(62)		183		(756)	(107)		(47)	
Amounts reclassified from accumulated other												, ,	_			
comprehensive income						63		33		(101)			 5		(50)	
Net current period comprehensive income																
(loss)		821		317		63		(29)		82		(756)	(102)		(97)	
Balance at June 30	\$	(3,996)	\$	(4,512)	\$	(2,215)	\$	(1,987)	\$	13	\$	(691)	\$ (52)	\$	(33)	

Reclassified amounts for foreign currency translation are recorded in the Condensed Consolidated Statement of Earnings as Net foreign exchange loss (gain); gains (losses) on marketable equity securities as Other expense (income), net and cash flow hedges as Cost of products sold. Net actuarial losses and prior service cost are included as a component of net periodic benefit plan costs; see Note 13 for additional details.

On January 4, 2017, Abbott completed the acquisition of St. Jude Medical, a global medical device manufacturer, for approximately \$23.6 billion, including approximately \$13.6 billion in cash and approximately \$10 billion in Abbott common shares, which represented approximately 254 million shares of Abbott common stock, based on Abbott's closing stock price on the acquisition date. As part of the acquisition, approximately \$5.9 billion of St. Jude Medical's debt was assumed, repaid or refinanced by Abbott. The transaction provides expanded opportunities for future growth and is an important part of the company's ongoing effort to develop a strong, diverse portfolio of devices, diagnostics, nutritionals and branded generic pharmaceuticals. The combined company will compete in nearly every area of the cardiovascular market, as well as in the neuromodulation market.

Under the terms of the agreement, for each St. Jude Medical common share, St. Jude Medical shareholders received \$46.75 in cash and 0.8708 of an Abbott common share. At an Abbott stock price of \$39.36, which reflects the closing price on January 4, 2017, this represented a value of approximately \$81 per St. Jude Medical common share and total purchase consideration of \$23.6 billion. The cash portion of the acquisition was funded through a combination of medium and long-term debt issued in November 2016 and a \$2.0 billion 120-day senior unsecured bridge term loan facility which was subsequently repaid.

The preliminary allocation of the fair value of the St. Jude Medical acquisition is shown in the table below. During the second quarter of 2017, measurement period adjustments to the value of the intangibles resulted in a credit of approximately \$70 million to intangible amortization expense to reduce the expense recorded in the first quarter of 2017. The allocation of the fair value of the acquisition will be finalized when the valuation is completed and differences between the preliminary and final allocation could be material.

(in billions)	
Acquired intangible assets, non-deductible	\$ 15.0
Goodwill, non-deductible	15.1
Acquired net tangible assets	3.4
Deferred income taxes recorded at acquisition	(4.6)
Net debt	(5.3)
Total preliminary allocation of fair value	\$ 23.6

The goodwill is primarily attributable to expected synergies from combining operations as well as intangible assets that do not qualify for separate recognition. The acquired tangible assets consist primarily of trade accounts receivable of approximately \$1.2 billion, inventory of approximately \$1.7 billion, other current assets of \$206 million, property and equipment of approximately \$1.5 billion, and other long-term assets of \$475 million. The acquired tangible liabilities consist of trade accounts payable and other current liabilities of approximately \$1.0 billion and other non-current liabilities of approximately \$655 million

If the acquisition of St. Jude Medical had occurred at the beginning of 2016, unaudited pro forma consolidated net sales would have been approximately \$13.2 billion and unaudited pro forma consolidated net loss would have been approximately \$165 million for the first six months of 2016, which includes the amortization of approximately \$820 million of inventory step-up and \$670 million of intangibles related to St. Jude Medical. For the second quarter of 2016, unaudited pro forma consolidated net sales would have been approximately \$6.9 billion and unaudited pro forma consolidated net earnings would have been approximately \$250 million, which includes the amortization of approximately \$430 million of inventory step-up and \$340 million of intangibles related to St. Jude Medical. The unaudited pro forma information is not necessarily indicative of the consolidated results of operations that would have been realized had the St. Jude Medical acquisition been completed as of the beginning of 2016, nor is it meant to be indicative of future results of operations that the combined entity will experience.

In the first six months of 2017, consolidated Abbott results include \$2.9 billion of sales and a pre-tax loss of approximately \$1.0 billion related to the St. Jude Medical acquisition, including approximately \$670 million of intangible amortization and \$820 million of inventory step-up amortization. It excludes acquisition, integration and restructuring-related costs.

In 2016, Abbott and St. Jude Medical agreed to sell certain businesses to Terumo Corporation for approximately \$1.12 billion. The sale included the St. Jude Medical Angio-SealTM and FemosealTM vascular closure and Abbott's Vado® Steerable Sheath businesses. The sale closed on January 20, 2017 and no gain or loss was recorded in the Condensed Consolidated Statement of Earnings.

On January 30, 2016, Abbott entered into a definitive agreement to acquire Alere Inc. (Alere), a diagnostic device and service provider, for \$56.00 per common share in cash. On April 13, 2017, Abbott and Alere amended the terms of the agreement to reduce the purchase price to \$51.00 per common share. The amended terms reduce the originally expected equity value by approximately \$500 million to a new expected equity value of approximately \$5.3 billion, which includes both common and preferred shares. On

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July 7, 2017, the Alere shareholders approved the acquisition. The acquisition is expected to close by the end of the third quarter of 2017, subject to the satisfaction of customary closing conditions, including applicable regulatory approvals. Under the amended terms of the acquisition agreement, the date by which necessary regulatory approvals must be received has been extended to September 30, 2017. The companies also agreed to dismiss their respective lawsuits. The acquisition is expected to significantly expand Abbott's global diagnostics presence and leadership. Abbott expects to utilize a combination of cash on hand and debt to fund the acquisition. Alere's net debt, which totaled \$2.3 billion at March 31, 2017, will be assumed, refinanced or repaid by Abbott.

On July 15, 2017, Alere entered into an agreement to sell its Triage MeterPro cardiovascular and toxicology business and the assets and liabilities related to its B-type Natriuretic Peptide assay business run on Beckman Coulter analyzers to Quidel Corporation (Quidel). The transactions with Quidel reflect a total purchase price of approximately \$400 million and contingent consideration with a maximum value of \$40 million. On July 21, 2017, Alere entered into an agreement with Siemens Diagnostics Holding II B.V. to sell its subsidiary, Epocal Inc., for approximately \$200 million. These transactions are subject to the successful completion of Abbott's acquisition of Alere and antitrust regulatory approvals. Alere is divesting these businesses in connection with the review by the Federal Trade Commission and the European Commission of Abbott's agreement to acquire Alere.

On July 17, 2017, Abbott commenced a tender offer to purchase for cash the 1.77 million outstanding shares of Alere's Series B Convertible Perpetual Preferred Stock at a price of \$402 per share, plus accrued but unpaid dividends to, but not including, the settlement date of the tender offer. This tender offer is subject to the satisfaction of certain conditions, including Abbott's acquisition of Alere and upon there being validly tendered (and not properly withdrawn)

at the expiration date of the tender offer that number of shares of Preferred Stock that equals at least a majority of the Preferred Stock issued and outstanding at the expiration of the tender offer.

On July 31, 2017, Abbott entered into a term loan agreement whereby Abbott can borrow up to \$2.8 billion on an unsecured basis for the acquisition of Alere. Funding of the 5-year loan is subject to the satisfaction of certain conditions related to the consummation of the Alere acquisition. Borrowings under the term loan will bear interest, at Abbott's option, based on either a base rate or a Eurodollar rate, plus an applicable margin based on Abbott's credit ratings.

Note 7 — Goodwill and Intangible Assets

The total amount of goodwill reported was \$22.132 billion at June 30, 2017 and \$7.683 billion at December 31, 2016. Goodwill increased by \$15.1 billion during the first six months of 2017 due to the completion of the St. Jude Medical acquisition, partially offset by a decrease of \$1.1 billion due to the sale of certain businesses to Terumo Corporation. Foreign currency translation adjustments increased goodwill by approximately \$421 million in the first six months of 2017. The amount reported at December 31, 2016 excludes goodwill reported in non-current assets held for disposition. As part of the sale of AMO in the first six months of 2017, approximately \$2.0 billion of goodwill was included as part of the net assets sold. The amount of goodwill related to reportable segments at June 30, 2017 was \$3.1 billion for the Established Pharmaceutical Products segment, \$286 million for the Nutritional Products segment, \$417 million for the Diagnostic Products segment, and \$17.3 billion for the Cardiovascular and Neuromodulation Products segment. The Cardiovascular and Neuromodulation Products segment as well as the goodwill related to the St. Jude Medical acquisition. There was no significant reduction of goodwill relating to impairments.

The gross amount of amortizable intangible assets, primarily product rights and technology was \$21.7 billion as of June 30, 2017 and \$10.4 billion as of December 31, 2016, and accumulated amortization was \$7.0 billion as of June 30, 2017 and \$6.2 billion as of December 31, 2016. The gross amount of amortizable intangible assets increased by \$11.3 billion during the first six months of 2017 due to the completion of the St. Jude Medical acquisition. Foreign currency translation adjustments increased intangible assets by \$128 million during the first six months of 2017. The December 31, 2016 amounts exclude net intangible assets reported in non-current assets held for disposition. As part of the sale of AMO in the first six months of 2017, approximately \$529 million of net intangible assets were included in the net assets sold.

Indefinite-lived intangible assets, which relate to in-process research and development acquired in a business combination, were approximately \$4.0 billion and \$349 million as of June 30, 2017 and December 31, 2016, respectively. Indefinite-lived intangible assets increased by \$3.7 billion due to the completion of the St. Jude Medical acquisition. In the first six months of 2016, Abbott recorded an impairment of a \$43 million in-process research and development project related to a non-reportable segment. Abbott's estimated annual amortization expense for intangible assets is approximately \$1.9 billion in 2017, \$2.1 billion in 2018, \$2.0 billion in 2019, \$1.8 billion in 2020 and \$1.7 billion in 2021. Amortizable intangible assets are amortized over 2 to 20 years (weighted average 11 years).

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Note 8 — Restructuring Plans

In 2017, Abbott management approved restructuring plans as part of the integration of the acquisition of St. Jude Medical into the cardiovascular and neuromodulation segment to leverage economies of scale and reduce costs. In the first six months of 2017, charges of approximately \$144 million, including one-time employee termination benefits were recorded as Selling, general and administrative expense. Abbott also assumed restructuring liabilities of approximately \$20 million as part of the St Jude Medical acquisition. The following summarizes the activity for the first six months of 2017 related to these actions and the status of the related accrual as of June 30, 2017:

(in millions)	_	
Liabilities assumed as part of business acquisition	\$	20
Restructuring charges recorded in 2017		144
Payments and other adjustments		(70)
Accrued balance at June 30, 2017	\$	94

From 2014 to 2017, Abbott management approved plans to streamline operations in order to reduce costs and improve efficiencies in various Abbott businesses including the nutritional and established pharmaceuticals businesses. In the first six months of 2017, charges of approximately \$17 million were recognized, of which approximately \$4 million is recorded in Cost of products sold, approximately \$7 million is recorded in Research and development and approximately \$6 million as Selling, general and administrative expense. The following summarizes the activity for the first six months of 2017 related to these restructuring actions and the status of the related accrual as of June 30, 2017:

(in millions)	
Accrued balance at December 31, 2016	\$ 66
Restructuring charges recorded in 2017	17
Payments and other adjustments	(22)
Accrued balance at June 30, 2017	\$ 61

Note 9 — Incentive Stock Programs

In connection with the completion of the St. Jude Medical acquisition in the first quarter of 2017, unvested St. Jude Medical stock options and restricted stock units were assumed by Abbott and converted into Abbott options and restricted stock units (as applicable) of substantially equivalent value, in accordance with the merger agreement. The number of shares underlying the converted options was 7,364,571 at a weighted average exercise price of \$30.50. The number of restricted stock units converted was 2,324,500 at a weighted average grant date fair value of \$37.69.

In the first six months of 2017, Abbott granted 4,370,403 stock options, 546,383 restricted stock awards and 6,865,354 restricted stock units under its incentive stock programs. At June 30, 2017, approximately 170 million shares were reserved for future grants. This reserve reflects the shares authorized by Abbott's shareholders in April 2017. Information regarding the number of options outstanding and exercisable at June 30, 2017 is as follows:

Outstanding	Exercisable

Number of shares	40,	577,408	23,873,390
Weighted average remaining life (years)		6.0	4.7
Weighted average exercise price	\$	35.72	\$ 33.92
Aggregate intrinsic value (in millions)	\$	523	\$ 351

The total unrecognized share-based compensation cost at June 30, 2017 amounted to approximately \$393 million which is expected to be recognized over the next three years.

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Note 10 — Debt and Lines of Credit

In the first six months of 2017, as part of the acquisition of St. Jude Medical, Abbott's long-term debt increased due to the assumption of outstanding debt previously issued by St. Jude Medical. Abbott exchanged certain St. Jude Medical debt obligations with an aggregate principal amount of approximately \$2.9 billion for debt issued by Abbott which consists of:

	Principal Amount
2.00% Senior Notes due 2018	\$473.8 million
2.80% Senior Notes due 2020	\$483.7 million
3.25% Senior Notes due 2023	\$818.4 million
3.875% Senior Notes due 2025	\$490.7 million
4.75% Senior Notes due 2043	\$639.1 million

Following this exchange, approximately \$194.2 million of existing St. Jude Medical notes remain outstanding across the five series of existing notes which have the same coupons and maturities as those listed above. There were no significant costs associated with the exchange of debt.

In addition, during the first six months of 2017, Abbott assumed and subsequently repaid the following St. Jude Medical debt obligations:

Term loan due 2020	\$2.3 billion
Yen-denominated notes due 2017 and 2020	\$179 million
Yen-denominated credit facilities	\$55 million
Commercial paper borrowings	\$220 million

On January 4, 2017, as part of funding the cash portion of the St. Jude Medical acquisition, Abbott borrowed \$2.0 billion under a 120-day senior unsecured bridge term loan facility. This facility was repaid during the first six months of 2017.

During the first six months of 2017, Abbott issued 364-day yen-denominated debt, of which \$195 million was outstanding at June 30, 2017. Abbott also paid off a \$479 million yen-denominated short-term debt.

In February 2016, Abbott obtained a commitment for a 364-day senior unsecured bridge term loan facility for an amount not to exceed \$9 billion in conjunction with its pending acquisition of Alere. This commitment, which was automatically extended for up to 90 days on January 29, 2017, expired on April 30, 2017 and was not renewed since Abbott does not need this bridge facility to finance the Alere acquisition. The fees associated with the bridge facilities were recognized in interest expense.

On July 31, 2017, Abbott entered into a term loan agreement whereby Abbott can borrow up to \$2.8 billion on an unsecured basis for the acquisition of Alere. Funding of the 5-year loan is subject to the satisfaction of certain conditions related to the consummation of the Alere acquisition. Borrowings under the term loan will bear interest, at Abbott's option, based on either a base rate or a Eurodollar rate, plus an applicable margin based on Abbott's credit ratings.

Note 11 — Financial Instruments, Derivatives and Fair Value Measures

Certain Abbott foreign subsidiaries enter into foreign currency forward exchange contracts to manage exposures to changes in foreign exchange rates for anticipated intercompany purchases by those subsidiaries whose functional currencies are not the U.S. dollar. These contracts, with gross notional amounts totaling \$2.8 billion at June 30, 2017 and \$2.6 billion at December 31, 2016 are designated as cash flow hedges of the variability of the cash flows due to changes in foreign exchange rates and are recorded at fair value. Accumulated gains and losses as of June 30, 2017 will be included in Cost of products sold at the time the products are sold, generally through the next twelve to eighteen months. The amount of hedge ineffectiveness was not significant in 2017 and 2016.

Abbott enters into foreign currency forward exchange contracts to manage currency exposures for foreign currency denominated third-party trade payables and receivables, and for intercompany loans and trade accounts payable where the receivable or payable is denominated in a currency other than the functional currency of the entity. For intercompany loans, the contracts require Abbott to sell or buy foreign currencies, primarily European currencies including the British pound, in exchange for primarily U.S. dollars and other European currencies. For intercompany and trade payables and receivables, the currency exposures are primarily the U.S. dollar and European currencies. At June 30, 2017 and December 31, 2016, Abbott held the gross notional amount of \$14.3 billion and \$14.9 billion, respectively, of such foreign currency forward exchange contracts.

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In March 2017, Abbott repaid its \$479 million foreign denominated short-term debt which was designated as a hedge of the net investment in a foreign subsidiary. At December 31, 2016, the value of this short-term debt was \$454 million and changes in the fair value of the debt up through the date of repayment due to changes in exchange rates were recorded in Accumulated other comprehensive income (loss), net of tax.

Abbott is a party to interest rate hedge contracts totaling approximately \$4.0 billion at June 30, 2017 and \$5.5 billion at December 31, 2016 to manage its exposure to changes in the fair value of fixed-rate debt. In the second quarter of 2017, Abbott unwound approximately \$1.5 billion in interest rate swaps relating to the 2.00% Note due in 2020 and the 2.55% Note due in 2022. The proceeds received were not significant. These contracts are designated as fair value hedges of the variability of the fair value of fixed-rate debt due to changes in the long-term benchmark interest rates. The effect of the hedge is to change a fixed-rate interest obligation to a variable rate for that portion of the debt. Abbott records the contracts at fair value and adjusts the carrying amount of the fixed-rate debt by an offsetting amount. The amount of hedge ineffectiveness was not significant in 2017 and 2016.

The following table summarizes the amounts and location of certain derivative financial instruments as of June 30, 2017 and December 31, 2016:

		Fair Val	lue - Assets	Fair Value - Liabilities									
(in millions)	June 30, 2017	Dec. 31, 2016	Balance Sheet Caption	June 30, 2017	Dec. 31, 2016	Balance Sheet Caption							
Interest rate swaps designated as fair value hedges	\$ 7	\$ 8	Deferred income taxes and other assets	\$ 59	\$ 74	Post-employment obligations, deferred income taxes and other long-term liabilities							
Foreign currency forward exchange													
contracts:													
Hedging instruments	29	99	Prepaid expenses and other receivables	85	15	Other accrued liabilities							
Others not designated as hedges	176	177	Prepaid expenses and other receivables	112	67	Other accrued liabilities							
Debt designated as a hedge of net investment in a foreign subsidiary	_	_	n/a	_	454	Short-term borrowings							
	\$ 212	\$ 284		\$ 256	\$ 610								

The following table summarizes the activity for foreign currency forward exchange contracts designated as cash flow hedges, debt designated as a hedge of net investment in a foreign subsidiary and certain other derivative financial instruments, as well as the amounts and location of income (expense) and gain (loss) reclassified into income for the three months and six months ended June 30, 2017 and 2016. The amount of hedge ineffectiveness was not significant in 2017 and 2016 for these hedges.

(in millions)	_	Gain (loss) Recogn Comprehensive I Three Months Ended June 30 2017 2016		E Inc			Income (expense Reclassified Three Months Ended June 30 2017 2016				nths une 3		Income Statement Caption			
Foreign currency forward exchange contracts designated as cash flow hedges	\$	(54)	\$	7	\$	(145)	\$	(51)	\$	(2)	\$	23	\$ (7)	\$	54	Cost of products sold
Debt designated as a hedge of net investment in a foreign subsidiary		_		(45)		(25)		(77)		n/a		n/a	n/a		n/a	n/a
Interest rate swaps designated as fair value hedges		n/a		n/a		n/a		n/a		40		44	14		145	Interest expense
								15								

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Losses of \$51 million and gains of \$20 million were recognized in the three months ended June 30, 2017 and 2016, respectively, related to foreign currency forward exchange contracts not designated as a hedge. Losses of \$42 million and gains of \$18 million were recognized in the six months ended June 30, 2017 and 2016, respectively, related to foreign currency forward exchange contracts not designated as a hedge. These amounts are reported in the Condensed Consolidated Statement of Earnings on the Net foreign exchange loss (gain) line.

The interest rate swaps are designated as fair value hedges of the variability of the fair value of fixed-rate debt due to changes in the long-term benchmark interest rates. The hedged debt is marked to market, offsetting the effect of marking the interest rate swaps to market.

The carrying values and fair values of certain financial instruments as of June 30, 2017 and December 31, 2016 are shown in the following table. The carrying values of all other financial instruments approximate their estimated fair values. The counterparties to financial instruments consist of select major international financial institutions. Abbott does not expect any losses from nonperformance by these counterparties.

December 31 2016

June 30 2017

		June 30	, 201	/	December 31, 2010						
(in millions)		Carrying Value		Fair Value	Carrying Value		Fair Value				
Investment Securities:											
Equity securities	\$	1,482	\$	1,482	\$ 2,906	\$	2,906				
Other		63		63	41		42				
Total Long-term Debt		(23,813)		(25,007)	(20,684)		(21,147)				
Foreign Currency Forward Exchange Contracts:											
Receivable position		205		205	276		276				
(Payable) position		(197)		(197)	(82)		(82)				
Interest Rate Hedge Contracts:											
Receivable position		7		7	8		8				

(Payable) position (59) (59) (74)

The fair value of the debt was determined based on significant other observable inputs, including current interest rates.

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The following table summarizes the bases used to measure certain assets and liabilities at fair value on a recurring basis in the balance sheet:

			Basis of Fair Value Measurement							
(in millions)	Outstanding Balances			Quoted Prices in Active Markets		Significant Other Observable Inputs	Significant Unobservable Inputs			
June 30, 2017:										
Equity securities	\$	1,183	\$	1,183	\$	_	\$			
Interest rate swap derivative financial instruments		7		_		7		_		
Foreign currency forward exchange contracts		205				205		<u> </u>		
Total Assets	\$	1,395	\$	1,183	\$	212	\$	_		
Fair value of hedged long-term debt	\$	3,938	\$	_	\$	3,938	\$	_		
Foreign currency forward exchange contracts		197		_		197		_		
Interest rate swap derivative financial instruments		59		_		59		_		
Contingent consideration related to business combinations		146		_		_		146		
Total Liabilities	\$	4,340	\$	_	\$	4,194	\$	146		
December 31, 2016:										
Equity securities	\$	2,676	\$	2,676	\$	_	\$	_		
Interest rate swap derivative financial instruments		8		_		8		_		
Foreign currency forward exchange contracts		276		_		276		_		
Total Assets	\$	2,960	\$	2,676	\$	284	\$	_		
Fair value of hedged long-term debt	\$	5,413	\$	_	\$	5,413	\$	_		
Interest rate swap derivative financial instruments		74		_		74		_		
Foreign currency forward exchange contracts		82		_		82		_		
Contingent consideration related to business combinations		136		_		_		136		
Total Liabilities	\$	5,705	\$	_	\$	5,569	\$	136		

Equity securities are principally comprised of Mylan N.V. ordinary shares. The fair value of the Mylan equity securities was determined based on the value of the publicly-traded ordinary shares. In the first six months of 2017, Abbott sold 50 million ordinary shares of Mylan N.V which had a value of \$1.9 billion. As a result of this sale, Abbott's ownership interest in Mylan N.V. decreased from approximately 14% to approximately 3.7%. The fair value of debt was determined based on the face value of the debt adjusted for the fair value of the interest rate swaps, which is based on a discounted cash flow analysis. The fair value of foreign currency forward exchange contracts is determined using a market approach, which utilizes values for comparable derivative instruments. The fair value of the contingent consideration was determined based on an independent appraisal adjusted for the time value of money and other changes in fair value. In the first six months of 2017, the increase in the fair value of the contingent consideration was due to the assumption of St. Jude Medical's contingent consideration obligations.

Note 12 — Litigation and Environmental Matters

Abbott has been identified as a potentially responsible party for investigation and cleanup costs at a number of locations in the United States and Puerto Rico under federal and state remediation laws and is investigating potential contamination at a number of company-owned locations. Abbott has recorded an estimated cleanup cost for each site for which management believes Abbott has a probable loss exposure. No individual site cleanup exposure is expected to exceed \$4 million, and the aggregate cleanup exposure is not expected to exceed \$10 million.

Abbott is involved in various claims and legal proceedings, and Abbott estimates the range of possible loss for its legal proceedings and environmental exposures to be from approximately \$40 million to \$45 million. The recorded accrual balance at June 30, 2017 for these proceedings and exposures was approximately \$45 million. This accrual represents management's best estimate of probable loss, as defined by FASB ASC No. 450, "Contingencies." Within the next year, legal proceedings may occur that may result in a change in the estimated loss accrued by Abbott. While it is not feasible to predict the outcome of all such proceedings and exposures with certainty, management believes that their ultimate disposition should not have a material adverse effect on Abbott's financial position, cash flows, or results of operations.

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Note 13 — Post-Employment Benefits

Retirement plans consist of defined benefit, defined contribution, and medical and dental plans. Net cost recognized in continuing operations for the three months and six months ended June 30 for Abbott's major defined benefit plans and post-employment medical and dental benefit plans is as follows:

_		Defined Ber	nefit Plans		Medical and Dental Plans				
	Three Months		Six Mo	nths	Three M	Ionths	Six Months		
	Ended June 30		Ended J	une 30	Ended J	une 30	Ended June 30		
(in millions)	2017	2016	2017	2016	2017	2016	2017	2016	

Service cost - benefits earned during the period	\$ 70	\$ 67	\$ 142 \$	134	\$ 6 \$	6 \$	13 \$	13
Interest cost on projected benefit								
obligations	71	73	143	146	11	10	22	22
Expected return on plan assets	(153)	(143)	(305)	(284)	(8)	(8)	(16)	(17)
Net amortization of:								
Actuarial loss, net	40	31	82	63	6	3	12	9
Prior service cost (credit)	_	_	_	_	(12)	(11)	(23)	(22)
Net cost - continuing operations	\$ 28	\$ 28	\$ 62 \$	59	\$ 3 \$	 \$	8 \$	5

In the first six months of 2017, Abbott recognized a \$10 million curtailment gain related to the disposition of AMO.

Abbott funds its domestic defined benefit plans according to IRS funding limitations. International pension plans are funded according to similar regulations. In the first six months of 2017 and 2016, \$321 million and \$524 million, respectively, were contributed to defined benefit plans and \$11 million and \$9 million, respectively, were contributed to the post-employment medical and dental benefit plans.

Note 14 — Taxes on Earnings

Taxes on earnings from continuing operations reflect the estimated annual effective rates and include charges for interest and penalties. In the first six months of 2017, taxes on earnings from continuing operations include \$430 million of tax expense related to the gain on the sale of the AMO business, which is taxed at a discrete tax rate. Earnings from discontinued operations, net of tax, of \$46 million for the first six months of 2017 primarily reflects the recognition of net tax benefits as a result of the resolution of various tax positions related to prior years. In the first six months of 2016, taxes on earnings from continuing operations includes the impact of a net tax benefit of approximately \$145 million as a result of the resolution of various tax positions from prior years, partially offset by the unfavorable impact of non-deductible foreign exchange losses related to Venezuela. Earnings from discontinued operations, net of tax, in the first six months of 2016 reflects the recognition of \$266 million of net tax benefits primarily as a result of the resolution of various tax positions related to prior years. The conclusion of these tax matters decreased the gross amount of unrecognized tax benefits by approximately \$444 million.

Tax authorities in various jurisdictions regularly review Abbott's income tax filings. Abbott believes that it is reasonably possible that the recorded amount of gross unrecognized tax benefits may decrease between \$200 million and \$350 million, including cash adjustments, within the next twelve months as a result of concluding various domestic and international tax matters. In the U.S., Abbott's federal income tax returns are settled through 2013 and St. Jude Medical's federal income tax returns are settled through 2013 except for one item.

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Note 15 — Segment Information

Abbott's principal business is the discovery, development, manufacture and sale of a broad line of health care products. Abbott's products are generally sold directly to retailers, wholesalers, hospitals, health care facilities, laboratories, physicians' offices and government agencies throughout the world. On January 4, 2017, Abbott completed the acquisition of St. Jude Medical. Beginning with the first quarter of 2017, Abbott's cardiovascular and neuromodulation business includes the results of its historical Vascular Products segment and the results of the businesses acquired from St. Jude Medical from the date of acquisition.

Abbott's reportable segments are as follows:

Established Pharmaceutical Products — International sales of a broad line of branded generic pharmaceutical products.

Nutritional Products — Worldwide sales of a broad line of adult and pediatric nutritional products.

Diagnostic Products — Worldwide sales of diagnostic systems and tests for blood banks, hospitals, commercial laboratories and alternate-care testing sites. For segment reporting purposes, the Core Laboratories Diagnostics, Molecular Diagnostics, Point of Care and Ibis diagnostic divisions are aggregated and reported as the Diagnostic Products segment.

Cardiovascular and Neuromodulation Products — Worldwide sales of rhythm management, electrophysiology, heart failure, vascular, structural heart and neuromodulation products.

Non-reportable segments include AMO through the date of sale and Diabetes Care.

Abbott's underlying accounting records are maintained on a legal entity basis for government and public reporting requirements. Segment disclosures are on a performance basis consistent with internal management reporting. Intersegment transfers of inventory are recorded at standard cost and are not a measure of segment operating earnings. The cost of some corporate functions and the cost of certain employee benefits are charged to segments at predetermined rates that approximate cost. Remaining costs, if any, are not allocated to segments. In addition, intangible asset amortization is not allocated to operating segments, and intangible assets and goodwill are not included in the measure of each segment's assets. As a result of the acquisition of St. Jude Medical, the total assets of the Cardiovascular and Neuromodulation segment increased from \$1.425 billion at December 31, 2016 to \$5.250 billion at June 30, 2017. The following segment information has been prepared in accordance with the internal accounting policies of Abbott, as described above, and is not presented in accordance with generally accepted accounting principles applied to the consolidated financial statements.

	 Net Sales to External Customers						Operating Earnings								
	 Three Months Ended June 30									ee Months led June 30			Six Months Ended June 30		
(in millions)	2017		2016		2017		2016		2017		2016		2017		2016
Established Pharmaceutical Products	\$ 1,021	\$	980	\$	1,971	\$	1,868	\$	180	\$	192	\$	320	\$	340
Nutritional Products	1,731		1,740		3,373		3,411		392		368		743		710
Diagnostic Products	1,273		1,226		2,431		2,344		338		288		622		555
Cardiovascular and Neuromodulation Products (a)	2,260		782		4,363		1,467		689		290		1,308		537

Total Reportable Segments		6,285	4,728	12,138	9,09	0	1,599	1,138	2,993	2,142
Other		352	605	834	1,12	8				
Net Sales	\$	6,637 \$	5,333	\$ 12,972	\$ 10,21	8				
Corporate functions and benefit plans costs	_						(104)	(94)	(197)	(175)
Non-reportable segments							74	52	120	50
Net interest expense							(183)	(83)	(387)	(108)
Share-based compensation (b)							(92)	(62)	(263)	(214)
Amortization of intangible assets							(392)	(145)	(914)	(289)
Other, net (c)							(607)	(91)	(321)	(689)
Earnings from continuing operations before taxes						\$	295 \$	715 \$	1,031 \$	717

- (a) Operating earnings for the first six months of 2017 include certain costs previously reflected in corporate functions during the first three months of 2017.
- (b) Approximately 50 percent of the annual net cost of share-based awards will typically be recognized in the first quarter due to the timing of the granting of share-based awards.
- (c) Other, net for the six months ended June 30, 2017, includes the gain on the sale of the AMO business. Other, net for the three and six months ended June 30, 2017, includes inventory step-up amortization, restructuring charges and integration costs associated with the acquisition of St. Jude Medical. Other, net for the six months ended June 30, 2016, includes the \$477 million foreign currency loss related to operations in Venezuela and the \$43 million impairment of an in-process research and development project related to a non-reportable segment.

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

Financial Review - Results of Operations

Abbott's revenues are derived primarily from the sale of a broad line of health care products under short-term receivable arrangements. Patent protection and licenses, technological and performance features, and inclusion of Abbott's products under a contract most impact which products are sold; price controls, competition and rebates most impact the net selling prices of products; and foreign currency translation impacts the measurement of net sales and costs. Abbott's primary products are nutritional products, branded generic pharmaceuticals, diagnostic testing products and cardiovascular and neuromodulation products.

The following table details sales by reportable segment for the three months and six months ended June 30. Percent changes are versus the prior year and are based on unrounded numbers.

	Net Sales to External Customers										
(in millions)	I Ju	ee Months Ended une 30, 2017		ee Months Ended June 30, 2016	Total Change	Impact of Foreign Exchange	Total Change Excl. Foreign Exchange				
Established Pharmaceutical Products	\$	1,021	\$	980	4.1%	0.6%	3.5%				
Nutritional Products		1,731		1,740	(0.6)	(1.1)	0.5				
Diagnostic Products		1,273		1,226	3.8	(1.6)	5.4				
Cardiovascular and Neuromodulation											
Products		2,260		782	189.0	(1.1)	190.1				
Total Reportable Segments		6,285		4,728	32.9	(0.8)	33.7				
Other		352		605	(41.8)	(1.3)	(40.5)				
Net Sales	\$	6,637	\$	5,333	24.4	(0.9)	25.3				
Total U.S.	\$	2,360	\$	1,655	42.5	_	42.5				
			-	.							
Total International	\$	4,277	\$	3,678	16.3	(1.3)	17.6				

Net Sales to External Customers											
	Ended		Ended	Total Change	Impact of Foreign Exchange	Total Change Excl. Foreign Exchange					
\$	1,971	\$	1,868	5.5%	1.0%	4.5%					
	3,373		3,411	(1.1)	(0.8)	(0.3)					
	2,431		2,344	3.7	(1.4)	5.1					
	4,363		1,467	197.4	(1.1)	198.5					
	12,138		9,090	33.5	(0.7)	34.2					
	834		1,128	(26.0)	(1.3)	(24.7)					
\$	12,972	\$	10,218	27.0	(0.7)	27.7					
· ·	_										
\$	4,684	\$	3,186	47.0	_	47.0					
	\$	\$ 1,971 3,373 2,431 4,363 12,138 834 \$ 12,972	Ended June 30, 2017 \$ 1,971 \$ 3,373	Six Months Ended June 30, 2017 Six Months Ended June 30, 2016 \$ 1,971 \$ 1,868 3,373 3,411 2,431 2,344 4,363 1,467 12,138 9,090 834 1,128 \$ 12,972 \$ 10,218	Six Months Ended June 30, 2017 Six Months Ended June 30, 2016 Total Change \$ 1,971 \$ 1,868 5.5% 3,373 3,411 (1.1) 2,431 2,344 3.7 4,363 1,467 197.4 12,138 9,090 33.5 834 1,128 (26.0) \$ 12,972 \$ 10,218 27.0	Six Months Ended June 30, 2017 Six Months Ended June 30, 2016 Total Change Impact of Foreign Exchange \$ 1,971 \$ 1,868 5.5% 1.0% 3,373 3,411 (1.1) (0.8) 2,431 2,344 3.7 (1.4) 4,363 1,467 197.4 (1.1) 12,138 9,090 33.5 (0.7) 834 1,128 (26.0) (1.3) \$ 12,972 \$ 10,218 27.0 (0.7)					

Total International \$ 8,288 \$ 7,032 17.9 (1.0) 18.9

Note: In order to compute results excluding the impact of exchange rates, current year U.S. dollar sales are multiplied or divided, as appropriate, by the current year average foreign exchange rates and then those amounts are multiplied or divided, as appropriate, by the prior year average foreign exchange rates

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Net sales growth in the second quarter and first six months of 2017 was driven by the acquisition of St. Jude Medical, Inc. (St. Jude Medical) which was completed on January 4, 2017 as well as growth in the Established Pharmaceutical Products and Diagnostic Products segments excluding the impact of foreign exchange. Beginning in the first quarter of 2017, Abbott's cardiovascular and neuromodulation business included the results of its historical Vascular Products segment and the results of the businesses acquired from St. Jude Medical from the date of acquisition. The decrease in the Other category for the second quarter and first six months of 2017 reflects the sale of the Abbott Medical Optics (AMO) segment to Johnson & Johnson, partially offset by double-digit growth in Abbott's Diabetes Care business. The AMO segment was included in Abbott's results as a non-reportable segment up to February 27, 2017. Excluding the St. Jude Medical acquisition, AMO results and the impact of foreign exchange, total net sales increased 2.7 percent, U.S. sales increased 1.1 percent and international sales increased 3.4 percent in the second quarter of 2017. Excluding the St. Jude Medical acquisition, AMO results and the impact of foreign exchange, total net sales increased 3.0 percent, U.S. sales increased 3.0 percent and international sales increased 3.1 percent in the first six months of 2017

The table below provides detail by sales category for the six months ended June 30. Percent changes are versus the prior year and are based on unrounded numbers.

(in millions)	June 30, 2017		June 30, 2016		Total Change	Impact of Foreign Exchange	Total Change Excl. Foreign Exchange
Established Pharmaceutical Products —							
Key Emerging Markets	\$	1,528	\$	1,388	10.1%	1.9%	8.2%
Other Emerging Markets		443		480	(7.8)	(1.7)	(6.1)
Nutritionals —							
International Pediatric Nutritionals		1,023		1,111	(8.0)	(1.6)	(6.4)
U.S. Pediatric Nutritionals		891		828	7.7	_	7.7
International Adult Nutritionals		847		831	1.9	(1.4)	3.3
U.S. Adult Nutritionals		612		641	(4.5)	<u> </u>	(4.5)
Diagnostics —							
Core Laboratory		1,931		1,863	3.6	(1.7)	5.3
Molecular		226		227	(0.5)	(0.2)	(0.3)
Point of Care		274		254	8.0	(0.2)	8.2
Cardiovascular and Neuromodulation —							
Rhythm Management		1,063		_	n/m	n/m	n/m
Electrophysiology		659		7	n/m	n/m	n/m
Heart Failure		301		_	n/m	n/m	n/m
Vascular		1,434		1,287	11.4	(1.0)	12.4
Structural Heart		524		173	203.2	(1.9)	205.1
Neuromodulation		382		_	n/m	n/m	n/m

Key Emerging Markets for the Established Pharmaceutical Products business include India, Russia, Brazil and China, along with several other markets that represent the most attractive long-term growth opportunities for Abbott's branded generics product portfolio. Excluding the effect of foreign exchange, sales in the Key Emerging Markets increased 8.2 percent compared to the first six months of 2016. Strong growth in Russia, China, and several countries across Latin America was partially offset by the impact associated with implementation of a new Goods and Services Tax (GST) system in India during the second quarter of 2017. Excluding the transitory effect of GST on purchasing patterns in India, sales in Key Emerging Markets would have grown double-digits in the first half of 2017. The 6.1 percent decrease in Other Emerging Markets in the first six months of 2017 primarily reflects the unfavorable impact of Venezuelan operations. Excluding Venezuela and the effect of foreign exchange, sales in Other Emerging Markets increased 4.1 percent versus the first six months of 2016.

Excluding the effect of foreign exchange, International Pediatric Nutritional sales decreased 6.4 percent in the first half of 2017. Challenging conditions in the Chinese infant formula market continued to impact international performance. In the U.S., above-market Pediatric sales growth of 7.7 percent was led by the continued momentum of several recently launched infant formula products as well as growth of the *PediaSure*® toddler brand. Excluding the effect of foreign exchange, International Adult Nutritional sales increased 3.3 percent compared to the first half of 2016 led by continued market growth across priority geographies while U.S. Adult Nutritional sales decreased 4.5 percent due to competitive and market dynamics.

Excluding the effect of foreign exchange, the 5.1 percent increase in Diagnostics sales was primarily driven by share gains in the Core Laboratory and Point of Care markets in the U.S. and higher sales to various international markets. During the quarter, Abbott announced CE Mark of its new Alinity hq hematology system to identify and quantify different types of blood cells to help diagnose blood-related diseases.

Excluding the effect of foreign exchange, the 198.5 percent increase in the Cardiovascular and Neuromodulation Products segment was driven by the acquisition of St. Jude Medical which was completed on January 4, 2017. Excluding the impact of the acquisition as well as the impact of foreign exchange, sales in the Cardiovascular and Neuromodulation Products segment decreased 1.5 percent in the first six months of 2017 as lower coronary stent sales and the comparison impact from the favorable 2016 resolution of a third-party royalty agreement were partially offset by higher Structural Heart and endovascular sales

The gross profit margin percentage was 46.3 percent for the second quarter of 2017 compared to 54.4 percent for the second quarter of 2016. The gross profit margin percentage was 45.0 percent for the first six months of 2017 compared to 53.8 percent for the first six months of 2016. The decrease primarily reflects higher intangible amortization expense and inventory step-up amortization related to the St. Jude Medical acquisition.

Research and development expenses increased by \$165 million, or 47.5 percent, in the second quarter of 2017, and increased by \$333 million, or 45.9 percent, in the first six months of 2017, due primarily to the addition of the acquired St. Jude Medical business. For the six months ended June 30, 2017, research and development expenditures totaled \$482 million for the Cardiovascular and Neuromodulation Products segment, \$262 million for the Diagnostic Products segment, \$80 million for the Established Pharmaceutical Products segment and \$97 million for the Nutritional Products segment.

Selling, general and administrative expenses for the second quarter and first six months of 2017 increased 22.7 percent and 32.6 percent, respectively, due primarily to the addition of the acquired St. Jude Medical business as well as the incremental expenses to integrate St. Jude Medical with Abbott's existing vascular business, partially offset by the impact of cost improvement initiatives across various Abbott functions and businesses.

In April 2017, Abbott received a warning letter from the U.S. Food and Drug Administration (FDA) related to its manufacturing facility in Sylmar, CA which was acquired by Abbott on January 4, 2017 as part of the acquisition of St. Jude Medical. This facility manufactures implantable cardioverter defibrillators, cardiac resynchronization therapy defibrillators, and monitors. The warning letter relates to the FDA's observations from an inspection of this facility. Abbott has prepared a comprehensive plan of corrective actions and execution of the plan is progressing.

Business Acquisitions

On January 4, 2017, Abbott completed the acquisition of St. Jude Medical, a global medical device manufacturer, for approximately \$23.6 billion, including approximately \$13.6 billion in cash and approximately \$10 billion in Abbott common shares, which represented approximately 254 million shares of Abbott common stock, based on Abbott's closing stock price on the acquisition date. As part of the acquisition, approximately \$5.9 billion of St. Jude Medical's debt was assumed, repaid or refinanced by Abbott. The transaction provides expanded opportunities for future growth and is an important part of the company's ongoing effort to develop a strong, diverse portfolio of devices, diagnostics, nutritionals and branded generic pharmaceuticals. The combined company will compete in nearly every area of the cardiovascular market, as well as in the neuromodulation market.

Under the terms of the agreement, for each St. Jude Medical common share, St. Jude Medical shareholders received \$46.75 in cash and 0.8708 of an Abbott common share. At an Abbott stock price of \$39.36, which reflects the closing price on January 4, 2017, this represented a value of approximately \$81 per St. Jude Medical common share and total purchase consideration of \$23.6 billion. The cash portion of the acquisition was funded through a combination of medium and long-term debt issued in November 2016 and a \$2.0 billion 120-day senior unsecured bridge term loan facility which was subsequently repaid.

The preliminary allocation of the fair value of the St. Jude Medical acquisition is shown in the table below. During the second quarter of 2017, measurement period adjustments to the value of the intangibles resulted in a credit of approximately \$70 million to intangible amortization expense to reduce the expense recorded in the first quarter of 2017. The allocation of the fair value of the acquisition will be finalized when the valuation is completed and differences between the preliminary and final allocation could be material.

(in billions)		
Acquired intangible assets, non-deductible	\$	15.0
Goodwill, non-deductible		15.1
Acquired net tangible assets		3.4
Deferred income taxes recorded at acquisition		(4.6)
Net debt		(5.3)
Total preliminary allocation of fair value	\$	23.6
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The goodwill is primarily attributable to expected synergies from combining operations as well as intangible assets that do not qualify for separate recognition. The acquired tangible assets consist primarily of trade accounts receivable of approximately \$1.2 billion, inventory of approximately \$1.7 billion, other current assets of \$206 million, property and equipment of approximately \$1.5 billion, and other long-term assets of \$475 million. The acquired tangible liabilities consist of trade accounts payable and other current liabilities of approximately \$1.0 billion and other non-current liabilities of approximately \$655 million

If the acquisition of St. Jude Medical had occurred at the beginning of 2016, unaudited pro forma consolidated net sales would have been approximately \$13.2 billion and unaudited pro forma consolidated net loss would have been approximately \$165 million for the first six months of 2016, which includes the amortization of approximately \$820 million of inventory step-up and \$670 million of intangibles related to St. Jude Medical. For the second quarter of 2016, unaudited pro forma consolidated net sales would have been approximately \$6.9 billion and unaudited pro forma consolidated net earnings would have been approximately \$250 million, which includes the amortization of approximately \$430 million of inventory step-up and \$340 million of intangibles related to St. Jude Medical. The unaudited pro forma information is not necessarily indicative of the consolidated results of operations that would have been realized had the St. Jude Medical acquisition been completed as of the beginning of 2016, nor is it meant to be indicative of future results of operations that the combined entity will experience.

In the first six months of 2017, consolidated Abbott results include \$2.9 billion of sales and a pre-tax loss of approximately \$1.0 billion related to the St. Jude Medical acquisition, including approximately \$670 million of intangible amortization and \$820 million of inventory step-up amortization. It excludes acquisition, integration and restructuring-related costs.

In 2016, Abbott and St. Jude Medical agreed to sell certain businesses to Terumo Corporation for approximately \$1.12 billion. The sale included the St. Jude Medical Angio-SealTM and FemosealTM vascular closure and Abbott's Vado® Steerable Sheath businesses. The sale closed on January 20, 2017 and no gain or loss was recorded in the Condensed Consolidated Statement of Earnings.

On January 30, 2016, Abbott entered into a definitive agreement to acquire Alere Inc. (Alere), a diagnostic device and service provider, for \$56.00 per common share in cash. On April 13, 2017, Abbott and Alere amended the terms of the agreement to reduce the purchase price to \$51.00 per common share. The amended terms reduce the originally expected equity value by approximately \$500 million to a new expected equity value of approximately \$5.3 billion, which includes both common and preferred shares. On July 7, 2017, the Alere shareholders approved the acquisition. The acquisition is expected to close by the end of the third quarter of 2017, subject to the satisfaction of customary closing conditions, including applicable regulatory approvals. Under the amended terms of the acquisition agreement, the date by which necessary regulatory approvals must be received has been extended to September 30, 2017. The companies also agreed to dismiss their respective lawsuits. The acquisition is expected to significantly expand Abbott's global diagnostics presence and leadership. Abbott expects to utilize a combination of cash on hand and debt to fund the acquisition. Alere's net debt, which totaled \$2.3 billion at March 31, 2017, will be assumed, refinanced or repaid by Abbott.

On July 15, 2017, Alere entered into an agreement to sell its Triage MeterPro cardiovascular and toxicology business and the assets and liabilities related to its B-type Natriuretic Peptide assay business run on Beckman Coulter analyzers to Quidel Corporation (Quidel). The transactions with Quidel reflect a total purchase price of approximately \$400 million and contingent consideration with a maximum value of \$40 million. On July 21, 2017, Alere entered into an agreement with Siemens Diagnostics Holding II B.V. to sell its subsidiary, Epocal Inc., for approximately \$200 million. These transactions are subject to the successful completion of Abbott's acquisition of Alere and antitrust regulatory approvals. Alere is divesting these businesses in connection with the review by the Federal Trade Commission and the European Commission of Abbott's agreement to acquire Alere.

On July 17, 2017, Abbott commenced a tender offer to purchase for cash the 1.77 million outstanding shares of Alere's Series B Convertible Perpetual Preferred Stock at a price of \$402 per share, plus accrued but unpaid dividends to, but not including, the settlement date of the tender offer. This tender offer is subject to the satisfaction of certain conditions, including Abbott's acquisition of Alere and upon there being validly tendered (and not properly withdrawn) at the expiration date of the tender offer that number of shares of Preferred Stock that equals at least a majority of the Preferred Stock issued and outstanding at the expiration of the tender offer.

On July 31, 2017, Abbott entered into a term loan agreement whereby Abbott can borrow up to \$2.8 billion on an unsecured basis for the acquisition of Alere. Funding of the 5-year loan is subject to the satisfaction of certain conditions related to the consummation of the Alere acquisition. Borrowings under the term loan will bear interest, at Abbott's option, based on either a base rate or a Eurodollar rate, plus an applicable margin based on Abbott's credit ratings.

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Restructuring Plans

The results for the first six months of 2017 reflect charges under approved restructuring plans as part of the integration of the acquisition of St. Jude Medical as well as costs related to other actions associated with the company's plans to streamline various operations. Abbott recorded employee related severance and other charges of approximately \$161 million in the first six months of 2017 related to these initiatives. Approximately \$4 million is recognized in Cost of products sold, \$7 million is recognized in Research and development and approximately \$150 million is recognized in Selling, general and administrative expense. See Note 8 to the financial statements, "Restructuring Plans," for additional information regarding these charges.

Interest Expense (Income), net

Interest expense (income), net increased \$100 million in the second quarter of 2017 and \$279 million in the first six months of 2017 compared to 2016 due primarily to the \$15.1 billion of debt issued in November 2016 related to the financing of the St. Jude Medical acquisition which closed on January 4, 2017.

Taxes on Earnings from Continuing Operations

Taxes on earnings from continuing operations reflect the estimated annual effective rates and include charges for interest and penalties. In the first six months of 2017, taxes on earnings from continuing operations include \$430 million of tax expense related to the gain on the sale of the AMO business, which is taxed at a discrete tax rate. Earnings from discontinued operations, net of tax, of \$46 million for the first six months of 2017 primarily reflects the recognition of net tax benefits as a result of the resolution of various tax positions related to prior years. In the first six months of 2016, taxes on earnings from continuing operations includes the impact of a net tax benefit of approximately \$145 million as a result of the resolution of various tax positions from prior years, partially offset by the unfavorable impact of non-deductible foreign exchange losses related to Venezuela. Earnings from discontinued operations, net of tax, in the first six months of 2016 reflects the recognition of \$266 million of net tax benefits primarily as a result of the resolution of various tax positions related to prior years. The conclusion of these tax matters decreased the gross amount of unrecognized tax benefits by approximately \$444 million.

Tax authorities in various jurisdictions regularly review Abbott's income tax filings. Abbott believes that it is reasonably possible that the recorded amount of gross unrecognized tax benefits may decrease between \$200 million and \$350 million, including cash adjustments, within the next twelve months as a result of concluding various domestic and international tax matters. In the U.S., Abbott's federal income tax returns are settled through 2013 and St. Jude Medical's federal income tax returns are settled through 2013 except for one item.

Discontinued Operations

On February 27, 2015, Abbott completed the sale of its developed markets branded generics pharmaceuticals business to Mylan Inc. (Mylan) for 110 million shares (or approximately 22%) of a newly formed entity (Mylan N.V.) that combined Mylan's existing business and Abbott's developed markets branded generics pharmaceuticals business. Mylan N.V. is publicly traded. The shareholder agreement with Mylan N.V. includes voting and other restrictions that prevent Abbott from exercising significant influence over the operating and financial policies of Mylan N.V.

In April 2015, Abbott sold 40.25 million of its 110 million ordinary shares of Mylan N.V. As a result of this sale, Abbott's ownership interest in Mylan N.V. decreased to approximately 14%.

In March 2017, Abbott sold 44 million ordinary shares of Mylan N.V. and received approximately \$1.7 billion in proceeds from the sale of these shares. In June 2017, Abbott sold an additional 6 million ordinary shares of Mylan N.V. and received \$239 million in proceeds. Abbott recorded an immaterial pre-tax gain on the sale of these shares in the first six months of 2017. The gain was recognized in the Other expense (income), net line of the Condensed Consolidated Statement of Earnings. As a result of these share sales, Abbott's ownership interest in Mylan N.V. decreased from approximately 14% to approximately 3.7%.

On January 1, 2013, Abbott completed the separation of AbbVie Inc. (AbbVie), which was formed to hold Abbott's research-based proprietary pharmaceuticals business. Abbott has retained all liabilities for all U.S. federal and foreign income taxes on income prior to the separation, as well as certain non-income related taxes attributable to AbbVie's business prior to the separation. AbbVie generally will be liable for all other taxes attributable to its business.

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Earnings from discontinued operations, net of tax of \$46 million and \$260 million in the first six months of 2017 and 2016 were driven primarily by the recognition of net tax benefits as a result of the resolution of various tax positions related to AbbVie's operations for years prior to the separation.

In the first quarter of 2016, Abbott received an additional \$25 million of proceeds related to the expiration of a holdback agreement associated with the sale of the animal health business and reported an after-tax gain on the sale in discontinued operations of \$16 million.

Assets and Liabilities Held for Disposition

In September 2016, Abbott announced that it entered into a definitive agreement to sell Abbott Medical Optics (AMO), its vision care business, to Johnson & Johnson for \$4.325 billion in cash, subject to customary purchase price adjustments for cash, debt and working capital. The decision to sell AMO reflects Abbott's proactive shaping of its portfolio in line with its strategic priorities. In February 2017, Abbott completed the sale of AMO to Johnson & Johnson and recognized a pre-tax gain of \$1.151 billion, which was reported in the Other expense (income), net line of the Condensed Consolidated Statement of Earnings in the first six months of 2017. Abbott recorded an after-tax gain of \$721 million in the first six months of 2017 related to the sale of AMO.

The operating results of AMO up to the date of sale continued to be included in Earnings from Continuing Operations as they did not qualify for reporting as discontinued operations. For the three months ended June 30, 2017 and 2016, the AMO earnings before taxes included in Abbott's consolidated earnings were nil and \$13 million, respectively. For the first six months ended June 30, 2017 and 2016, the AMO losses before taxes included in Abbott's consolidated earnings were \$18 million and \$44 million, respectively. The following assets and liabilities of this business were reported as held for disposition in Abbott's Condensed Consolidated Balance Sheet as of December 31, 2016:

(in millions)	ember 31, 2016
Trade receivables, net	\$ 222
Total inventories	240
Prepaid expenses and other current assets	51
Current assets held for disposition	 513
Net property and equipment	247
Intangible assets, net of amortization	529
Goodwill	1,966
Deferred income taxes and other assets	11
Non-current assets held for disposition	2,753
Total assets held for disposition	\$ 3,266
Trade accounts payable	\$ 71
Salaries, wages, commissions and other accrued liabilities	174
Current liabilities held for disposition	 245
Post-employment obligations, deferred income taxes and other long-term liabilities	59
Total liabilities held for disposition	\$ 304

Liquidity and Capital Resources June 30, 2017 Compared with December 31, 2016

The reduction of cash and cash equivalents from \$18.6 billion at December 31, 2016 to \$9.7 billion at June 30, 2017 reflects the use of cash to fund the cash portion of the St. Jude Medical acquisition, repayments of debt, pension contributions, and dividends paid in the first six months of 2017, partially offset by proceeds from the disposition of businesses and sale of a portion of the Mylan ordinary shares.

Net cash from operating activities for the first six months of 2017 totaled \$1.925 billion, an increase of \$1.109 billion over the prior year due primarily to the favorable impact of the acquisition of the St. Jude Medical businesses, as well as a reduction in pension contributions. The Other, net line in Net cash from operating activities for the first six months of 2017 of \$50 million includes the impact of approximately \$430 million of tax expense associated with the disposition of businesses, which has not yet been paid. Other net, also includes contributions to defined benefit pension plans of \$321 million. The Other, net line in Net cash from operating activities for the first six months of 2016 of \$1.2 billion includes contributions to defined benefit pension plans of \$524 million as well as approximately \$140 million of cash taxes paid related to the disposition of businesses. Other, net in 2016 also includes the non-cash impact of approximately \$410 million of net tax benefits primarily associated with the resolution of various tax positions from prior

years. The foreign currency loss related to Venezuela in the first six months of 2016 reduced Abbott's cash by approximately \$410 million and is shown on the Effect of exchange rate changes on cash and cash equivalents line within the Condensed Consolidated Statement of Cash Flows. Abbott expects to fund cash dividends, capital expenditures and its other investments in its businesses with cash flow from operating activities, cash on hand, short-term investments and borrowings.

Working capital was \$13.2 billion at June 30, 2017 and \$20.1 billion at December 31, 2016. The \$6.9 billion decrease in working capital in 2017 is primarily due to the reduction in Cash and cash equivalents driven by the use of cash to fund the cash portion of the St. Jude Medical acquisition, debt repayments, pension contributions and dividend payments, partially offset by proceeds from the sale of a portion of the Mylan ordinary shares and business dispositions. In the first six months of 2017, Abbott sold 50 million ordinary shares of Mylan N.V. which generated cash proceeds of approximately \$1.9 billion.

Since January 2010, Venezuela has been designated as a highly inflationary economy under U.S. GAAP. In 2014 and 2015, the government of Venezuela operated multiple mechanisms to exchange bolivars into U.S. dollars. These mechanisms included the CENCOEX, SICAD, and SIMADI rates, which stood at 6.3, 13.5, and approximately 200, respectively, at December 31, 2015. In 2015, Abbott continued to use the CENCOEX rate of 6.3 Venezuelan bolivars to the U.S. dollar to report the results, financial position, and cash flows related to its operations in Venezuela since Abbott continued to qualify for this exchange rate to pay for the import of various products into Venezuela.

On February 17, 2016, the Venezuelan government announced that the three-tier exchange rate system would be reduced to two rates renamed the DIPRO and DICOM rates. The DIPRO rate is the official rate for food and medicine imports and was adjusted from 6.3 to 10 bolivars per U.S. dollar. The DICOM rate is a floating market rate published daily by the Venezuelan central bank, which at the end of the first quarter of 2016 was approximately 263 bolivars per U.S. dollar. As a result of decreasing government approvals to convert bolivars to U.S. dollars to pay for intercompany accounts, as well as the accelerating deterioration of economic conditions in the country, Abbott concluded that it was appropriate to move to the DICOM rate at the end of the first quarter of 2016. As a result, Abbott recorded a foreign currency exchange loss of \$477 million in the first quarter of 2016 to revalue its net monetary assets in Venezuela. Abbott is continuing to use the DICOM rate to report the results of operations and to remeasure net monetary assets for Venezuela at the end of each quarter. As of June 30, 2017, Abbott's Venezuelan operations represented approximately 0.01% of Abbott's consolidated assets and any additional foreign currency losses related to Venezuela are not expected to be material.

At June 30, 2017, Abbott's long-term debt rating was BBB by Standard & Poor's Corporation and Baa3 by Moody's Investors Service. Abbott expects to maintain an investment grade rating. Abbott has readily available financial resources, including unused lines of credit of \$5.0 billion which expire in 2019.

In September 2014, the board of directors authorized the repurchase of up to \$3.0 billion of Abbott's common shares from time to time. The 2014 authorization was in addition to the \$512 million unused portion of a previous program announced in June 2013. In the first six months of 2016, Abbott repurchased 10.4 million shares at a cost of \$408 million under the program authorized in 2014.

On April 27, 2016, the board of directors authorized the issuance and sale for general corporate purposes of up to 75 million common shares that would result in proceeds of up to \$3 billion. No shares have been issued under this authorization.

In each of the first two quarters of 2017, Abbott declared a quarterly dividend of \$0.265 per share on its common shares, which represents an increase of approximately 2% over the \$0.26 per share quarterly dividend declared in each of the first two quarters of 2016.

Recently Issued Accounting Standards

In March 2017, the Financial Accounting Standards Board (FASB) issued Accounting Standards Update (ASU) 2017-07, Compensation — Retirement Benefits (Topic 715): Improving the Presentation of Net Periodic Pension Cost and Net Periodic Postretirement Benefit Cost which changes the financial statement presentation requirements for pension and other postretirement benefit expense. While service cost will continue to be reported in the same financial statement line items as other current employee compensation costs, the ASU requires all other components of pension and other postretirement benefit expense to be presented separately from service cost, and outside any subtotal of income from operations. The standard becomes effective for Abbott beginning in the first quarter of 2018 and early adoption is permitted. Abbott is currently evaluating the impact ASU 2017-07 will have on its consolidated financial statements.

In October 2016, the FASB issued ASU 2016-16, *Income Taxes (Topic 740): Intra-Entity Transfers of Assets Other Than Inventory*, which requires the recognition of the income tax effects of intercompany sales and transfers of assets, other than inventory, in the period in which the transfer occurs. The standard becomes effective for Abbott beginning in the first quarter of 2018 and early adoption is permitted. Abbott is currently evaluating the impact ASU 2016-16 will have on its consolidated financial statements.

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In February 2016, the FASB issued ASU 2016-02, *Leases*, which requires lessees to recognize assets and liabilities for most leases on the balance sheet. The standard becomes effective for Abbott beginning in the first quarter of 2019 and early adoption is permitted. Adoption requires application of the new guidance for all periods presented. Abbott is currently evaluating the impact the new guidance will have on its consolidated financial statements.

In January 2016, the FASB issued ASU 2016-01, *Financial Instruments — Recognition and Measurement of Financial Assets and Financial Liabilities*, which provides new guidance for the recognition, measurement, presentation, and disclosure of financial assets and liabilities. The standard becomes effective for Abbott beginning in the first quarter of 2018 and early adoption is permitted. Abbott is currently evaluating the effect, if any, that the standard will have on its consolidated financial statements and related disclosures.

In May 2014, the FASB issued ASU 2014-09, *Revenue from Contracts with Customers*, which provides a single comprehensive model for accounting for revenue from contracts with customers and will supersede most existing revenue recognition guidance. The standard becomes effective for Abbott in the first quarter of 2018. Abbott's revenues are primarily comprised of product sales. Abbott has made substantial progress in the evaluation of the new standard including a detailed review of Abbott's revenue streams and contracts. Based on the work performed to date, Abbott currently does not expect the adoption of the new standard to have a material impact on its consolidated financial statements. Abbott is continuing to evaluate the effect that the standard will have on its consolidated financial statements including the new disclosure requirements. Abbott will continue to monitor additional modifications, clarifications or interpretations undertaken by the FASB that may impact Abbott's current conclusions. Abbott is currently expecting to use the modified retrospective method to adopt this standard.

Legislative Issues

Abbott's primary markets are highly competitive and subject to substantial government regulations throughout the world. Abbott expects debate to continue over the availability, method of delivery, and payment for health care products and services. It is not possible to predict the extent to which Abbott or the health care industry in general might be adversely affected by these factors in the future. A more complete discussion of these factors is contained in Item 1, Business, and Item 1A, Risk Factors, in the 2016 Annual Report on Form 10-K.

Private Securities Litigation Reform Act of 1995 — A Caution Concerning Forward-Looking Statements

Under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995, Abbott cautions investors that any forward-looking statements or projections made by Abbott, including those made in this document, are subject to risks and uncertainties that may cause actual results to differ materially from those projected. Economic, competitive, governmental, technological and other factors that may affect Abbott's operations are discussed in Item 1A, Risk Factors, in the 2016 Annual Report on Form 10-K.

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PART I. FINANCIAL INFORMATION

<u>Item 3.</u> <u>Quantitative and Qualitative Disclosures about Market Risk</u>

Market Price Sensitive Investments

The fair value of the available-for-sale equity securities held by Abbott was approximately \$1.2 billion as of June 30, 2017 and \$2.7 billion as of December 31, 2016. The decrease is due primarily to the sale of 50 million ordinary shares of Mylan N.V., thereby reducing Abbott's equity securities by approximately \$1.9 billion during the first six months of 2017. All available-for-sale equity securities are subject to potential changes in market value. A hypothetical 20 percent decrease in the share prices of these investments would decrease their fair value at June 30, 2017 by approximately \$240 million. Abbott monitors these investments for other than temporary declines in market value, and charges impairment losses to income when an other than temporary decline in value occurs.

Item 4. Controls and Procedures

- (a) Evaluation of disclosure controls and procedures. The Chief Executive Officer, Miles D. White, and Chief Financial Officer, Brian B. Yoor, evaluated the effectiveness of Abbott Laboratories' disclosure controls and procedures as of the end of the period covered by this report, and concluded that Abbott Laboratories' disclosure controls and procedures were effective to ensure that information Abbott is required to disclose in the reports that it files or submits with the Securities and Exchange Commission (the "Commission") under the Securities Exchange Act of 1934 (the "Exchange Act") is recorded, processed, summarized and reported, within the time periods specified in the Commission's rules and forms, and to ensure that information required to be disclosed by Abbott in the reports that it files or submits under the Exchange Act is accumulated and communicated to Abbott's management, including its principal executive officer and principal financial officer, as appropriate to allow timely decisions regarding required disclosure.
- (b) Changes in internal control over financial reporting. During the quarter ended June 30, 2017, there were no changes in Abbott's internal control over financial reporting (as defined in Rule 13a-15(f) under the Exchange Act) that have materially affected, or are reasonably likely to materially affect, Abbott's internal control over financial reporting.

PART II. OTHER INFORMATION

Item 1. Legal Proceedings

Abbott is involved in various claims, legal proceedings and investigations, including those described in our Annual Report on Form 10-K for the year ended December 31, 2016 and Quarterly Report on Form 10-Q for the quarterly period ended March 31, 2017.

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

(c) Issuer Purchases of Equity Securities

Period	(a) Total Number of Shares (or Units) Purchased	(b) Average Price Paid per Share (or Unit)	(c) Total Number of Shares (or Units) Purchased as Part of Publicly Announced Plans or Programs	Number (or Approximate Dollar Value) of Shares (or Units) that May Yet Be Purchased Under the Plans or Programs
April 1, 2017 — April 30, 2017	15,015(1)		_	\$ 925,131,209(2)
May 1, 2017 — May 31, 2017	17,515(1) \$	S 44.104	_	\$ 925,131,209(2)
June 1, 2017 — June 30, 2017	24,054(1) \$	47.600	_	\$ 925,131,209(2)
Total	56,584(1) \$	45.622	_	\$ 925,131,209(2)

(d) Maximum

- (1) These shares include:
 - (i) the shares deemed surrendered to Abbott to pay the exercise price in connection with the exercise of employee stock options 707 in April, 3,207 in May, and 9,746 in June; and
 - (ii) the shares purchased on the open market for the benefit of participants in the Abbott Laboratories, Limited Employee Stock Purchase Plan 14,308 in April, 14,308 in May, and 14,308 in June.

These shares do not include the shares surrendered to Abbott to satisfy tax withholding obligations in connection with the vesting of restricted stock or restricted stock units.

(2) On September 11, 2014, Abbott announced that its board of directors approved the purchase of up to \$3 billion of its common shares, from time to time.

Item 6. Exhibits

Incorporated by reference to the Exhibit Index included herewith.

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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.

ABBOTT LABORATORIES

By: /s/ Brian B. Yoor

Brian B. Yoor

Executive Vice President, Finance and Chief Financial Officer

Date: August 2, 2017

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EXHIBIT INDEX

Exhibit No.	<u>Exhibit</u>
2.1	Amendment to Agreement and Plan of Merger, dated as of April 13, 2017, among Alere Inc., Abbott Laboratories and Angel Sub, Inc., filed as Exhibit 2.1 to the Abbott Laboratories Current Report on Form 8-K dated April 14, 2017.
3.1	By-Laws of Abbott Laboratories, as amended and restated effective June 29, 2017, filed as Exhibit 3.1 to the Abbott Laboratories Current Report on Form 8-K dated June 29, 2017.
10.1	Abbott Laboratories 2017 Incentive Stock Program (incorporated by reference to Exhibit B of Abbott's Definitive Proxy Statement on Schedule 14A filed with the Securities and Exchange Commission on March 17, 2017).
10.2	Form of Restricted Stock Unit Agreement (ratably vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.2 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.3	Form of Restricted Stock Unit Agreement for foreign employees (ratably vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.3 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.4	Form of Restricted Stock Unit Agreement (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.4 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.5	Form of Restricted Stock Unit Agreement for foreign employees (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.5 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.6	Form of Performance Restricted Stock Unit Agreement for foreign employees (annual performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.6 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.7	Form of Performance Restricted Stock Unit Agreement for foreign employees (interim performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.7 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.8	Form of Restricted Stock Agreement (ratably vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.8 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.9	Form of Restricted Stock Agreement (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.9 to

	the Abbott Laboratories Current Report on Form 6-K dated April 26, 2017.
10.10	Form of Performance Restricted Stock Agreement (annual performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.10 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.11	Form of Performance Restricted Stock Agreement (interim performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.11 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.12	Form of Non-Qualified Stock Option Agreement under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.12 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.13	Form of Non-Qualified Stock Option Agreement for foreign employees under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.13 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.14	Form of Restricted Stock Unit Agreement for executive officers (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.14 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
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10.15	Form of Restricted Stock Unit Agreement for foreign executive officers (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.15 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.16	Form of Performance Restricted Stock Unit Agreement for foreign executive officers (annual performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.16 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.17	Form of Performance Restricted Stock Unit Agreement for foreign executive officers (interim performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.17 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.18	Form of Restricted Stock Agreement for executive officers (ratably vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.18 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.19	Form of Restricted Stock Agreement for executive officers (cliff vested) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.19 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.20	Form of Performance Restricted Stock Agreement for executive officers (annual performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.20 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.21	Form of Performance Restricted Stock Agreement for executive officers (interim performance based) under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.21 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.22	Form of Non-Qualified Stock Option Agreement for executive officers under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.22 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.23	Form of Non-Qualified Stock Option Agreement for foreign executive officers under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.23 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.24	Form of Non-Employee Director Restricted Stock Unit Agreement under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.24 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.25	Form of Non-Employee Director Restricted Stock Unit Agreement for foreign non-employee directors under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.25 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.26	Form of Non-Employee Director Non-Qualified Stock Option Agreement under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.26 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
10.27	Form of Non-Employee Director Non-Qualified Stock Option Agreement for foreign non-employee directors under the Abbott Laboratories 2017 Incentive Stock Program, filed as Exhibit 10.27 to the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.
12	Statement re: Computation of ratio of earnings to fixed charges.
31.1	Certification of Chief Executive Officer Required by Rule 13a-14(a) (17 CFR 240.13a-14(a)).
31.2	Certification of Chief Financial Officer Required by Rule 13a-14(a) (17 CFR 240.13a-14(a)).
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the Abbott Laboratories Current Report on Form 8-K dated April 28, 2017.

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- Certification of Chief Executive Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
- 32.2 Certification of Chief Financial Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
- The following financial statements and notes from the Abbott Laboratories Quarterly Report on Form 10-Q for the quarter and six months ended June 30, 2017, formatted in XBRL: (i) Condensed Consolidated Statement of Earnings; (ii) Condensed Consolidated Statement of Comprehensive Income; (iii) Condensed Consolidated Balance Sheet; (iv) Condensed Consolidated Statement of Cash Flows; and (v) Notes to Condensed Consolidated Financial Statements.

Abbott Laboratories and Subsidiaries

Computation of Ratio of Earnings to Fixed Charges

(Unaudited)

(dollars in millions)

	Six Months Ended June 30, 2017	
Earnings from Continuing Operations	\$	656
Add (deduct):		
Taxes on earnings		375
Capitalized interest cost, net of amortization		(6)
Noncontrolling interests		8
Earnings from Continuing Operations, as adjusted		1,033
Fixed Charges:		
Interest on long-term and short-term debt		440
Capitalized interest cost		14
Rental expense representative of an interest factor		57
		,
Total Fixed Charges		511
		,
Total adjusted earnings available for payment of fixed charges	\$	1,544
Ratio of earnings to fixed charges		3.0

NOTE: For the purpose of calculating this ratio, (i) earnings from continuing operations have been calculated by adjusting earnings for taxes on earnings; interest expense; capitalized interest cost, net of amortization; noncontrolling interests; and the portion of rentals representative of the interest factor, (ii) Abbott considers one-third of rental expense to be the amount representing return on capital, and (iii) fixed charges comprise total interest expense, including capitalized interest and such portion of rentals.

Certification of Chief Executive Officer Required by Rule 13a-14(a) (17 CFR 240.13a-14(a))

I, Miles D. White, certify that:

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

Date: August 2, 2017 /s/ Miles D. White

Miles D. White Chairman of the Board and Chief Executive Officer

Certification of Chief Financial Officer Required by Rule 13a-14(a) (17 CFR 240.13a-14(a))

I, Brian B. Yoor, certify that:

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

Date: August 2, 2017 /s/ Brian B. Yoor

Brian B. Yoor Executive Vice President, Finance and Chief Financial Officer

Certification Pursuant To 18 U.S.C. Section 1350 As Adopted Pursuant To Section 906 of the Sarbanes-Oxley Act of 2002

In connection with the Quarterly Report of Abbott Laboratories (the "Company") on Form 10-Q for the period ended June 30, 2017 as filed with the Securities and Exchange Commission (the "Report"), I, Miles D. White, Chairman of the Board and Chief Executive Officer of the Company, certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to § 906 of the Sarbanes-Oxley Act of 2002, that:

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

/s/ Miles D. White

Miles D. White Chairman of the Board and Chief Executive Officer August 2, 2017

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

Certification Pursuant To 18 U.S.C. Section 1350 As Adopted Pursuant To Section 906 of the Sarbanes-Oxley Act of 2002

In connection with the Quarterly Report of Abbott Laboratories (the "Company") on Form 10-Q for the period ended June 30, 2017 as filed with the Securities and Exchange Commission (the "Report"), I, Brian B. Yoor, Executive Vice President, Finance and Chief Financial Officer of the Company, certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to § 906 of the Sarbanes-Oxley Act of 2002, that:

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

/s/ Brian B. Yoor

Brian B. Yoor Executive Vice President, Finance and Chief Financial Officer August 2, 2017

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