



Supporting the heroes of healthcare

Annual Report 2017



BD associates have a passion and commitment to help improve patient outcomes, increase the safety and efficiency of clinicians' care delivery process, enable laboratory scientists to better diagnose disease and advance researchers' capabilities to develop the next generation of diagnostics and therapeutics.



Nearly

50,000

associates



Serving

190+

countries



Vincent A. Forlenza

Chairman and
Chief Executive Officer

To our shareholders, customers and associates:

In just a few short years,
we have transformed
BD like no other time in
our 120-year history.

We have accelerated our momentum in our core solutions, we have integrated CareFusion to enhance the value we offer to our customers, and we re-engineered our functional teams to leverage our scale and operate more efficiently and effectively. This groundwork will prepare the company for the next major milestone in our strategic transformation as we acquire and integrate C. R. Bard, Inc. I could not be more proud of our associates for the hard work they have exhibited every day to fulfill our purpose of *advancing the world of health* while continuing to deliver solid and dependable top- and bottom-line results.

As companies grow, they often become more complex. We have been deliberate in our approach to ensure that as we continue to grow both organically and inorganically, we continually simplify how we operate to reduce complexity and stay nimble. This approach is grounded in our deep focus and understanding of the role BD has in the healthcare ecosystem: **We help the people who help the patients.** That includes the caregivers, the laboratory professionals, the researchers and others in healthcare settings around the world that have direct responsibility for making patients better. They are our heroes.

We support these heroes of healthcare by providing solutions, services, education and expertise so that they can provide the best care possible and develop the next generation of diagnostics and therapies. We help the healers. And we don't take that responsibility lightly.

New solutions

The power of the "new" BD is seen in our new solutions. We no longer focus on just new products, but instead we determine how we can bring a suite of products and services to solve customer challenges. Everything starts with the customer.

We are redefining medication management technologies to support hospitals and health systems to make their process safer, smarter and simpler. Through the **BD HealthSight™** platform for enterprise medication management, we have introduced a new

There is a 45% chance for error in the medication administration process.

concept in the integration of technology, analytics and expert services to optimize the medication management process and help improve efficiency, reduce waste and improve patient outcomes.

About **90%** of hospital inpatients receive a peripheral IV, and those IVs have a **35% to 50%** failure rate.

We are also tackling IV-related complications associated with vascular access management through our ability to combine clinical practice assessments, a comprehensive portfolio of products, expert clinical training and education to help hospitals reduce unnecessary complications.

For clinical laboratories looking for better ways to diagnose, we offer a comprehensive set of tools to support disease diagnosis and treatment. This past year, we have introduced a wide variety of new ID/AST diagnostics that can identify infectious diseases and provide antimicrobial susceptibility data to enable physicians to prescribe appropriate treatments. Through **BD Kiestra™** lab automation solutions, we continue to automate clinical laboratories for maximum efficiency.

We are also on the front lines of helping medical researchers develop a new generation of diagnostics and therapies through our expertise in bioscience and genomics. The new **BD Rhapsody™** platform is a complete system of reagents, instruments and software for targeted gene expression in

single cell analysis with the ability to detect rare molecules that researchers use to study whether the presence of these biological markers are indicative of disease. As immunotherapy research continues to

Sequencing the genome: then and now

	2001	Today
Cost	\$2.7 billion	\$5,000
Time	13 years	<1 week

expand, the ability to isolate and characterize individual cells becomes exponentially more important, and BD is a recognized leader in cell sorting applications. This is an area where we will continue to lead.

And we continue to invest in innovation, including new technologies under development to provide more advanced “wearable” drug delivery devices and diabetes disease management applications.

The International Diabetes Federation forecasts that by 2040 over **600 million** people will suffer from diabetes, costing the healthcare system greater than **\$1 trillion**.

Leadership

Developing the next generation of solutions for clinicians and researchers is just the beginning of our industry leadership. As we grow into one of the top five largest medical technology companies in the world, we are also leading in areas that are critical to the future of our industry and to society at-large.

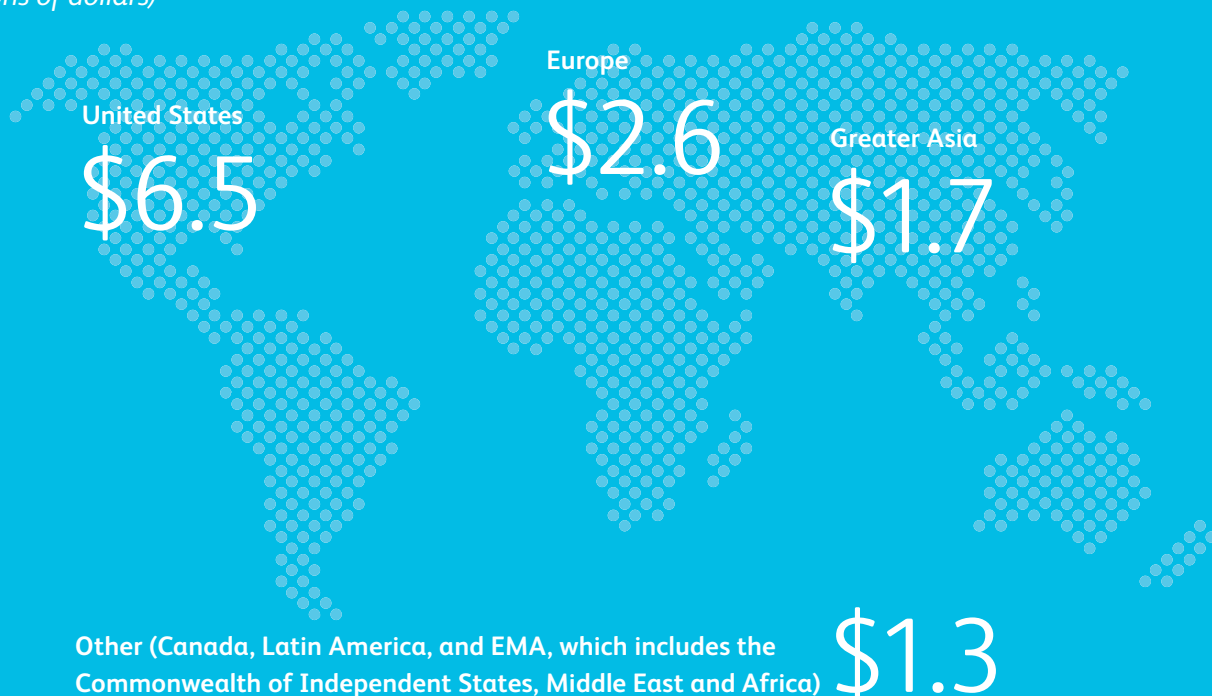
This is evident in our approach to emerging worldwide health concerns, such as antimicrobial resistance (AMR), which current trends project could cause up to 10 million

50% of antibiotic use in the U.S. is unnecessary.

deaths annually by 2050, surpassing deaths currently caused by cancer. To combat AMR, we have mobilized our internal teams across the company’s businesses, regions and functions to integrate our unique combination of expertise and solutions across infection prevention, diagnostics and surveillance. We are also engaging with external partners around the world through the new “**I’m a Resistance Fighter™**” AMR campaign, where BD is building a global coalition of organizations, clinicians and patients taking actionable steps to combat AMR.

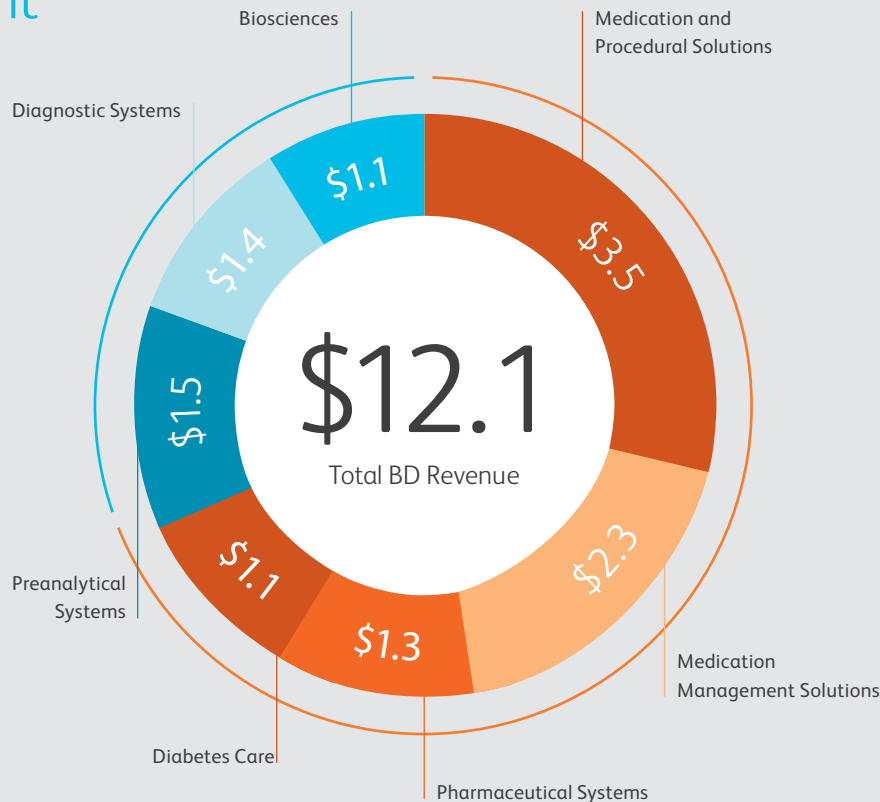
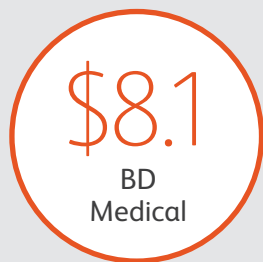
Revenue by geography

(billions of dollars)



Revenue by segment

(billions of dollars)



Values in this exhibit reflect rounded numbers.

We also take seriously the role we play as leaders in the communities where we live and work. BD committed up to \$800,000 in cash and/or product donations to be shared among our charitable partners to assist those impacted by the hurricanes, flooding and wildfires that affected the U.S. and Puerto Rico in 2017. We are humbled by the incredible perseverance and dedication our more than 600 associates in Puerto Rico demonstrated to restore our operations following Hurricane Maria, many working tirelessly to put our customers' and patients' needs ahead of their own.

Partnership and collaboration

Finding solutions to current and future challenges requires partnership and collaboration with other global leaders.

In 2017, the **BD Helping Build Healthy Communities™** partnership was selected as one of three winners for the Healthy10 Healthy Community award from the U.S. Chamber of Commerce Foundation. In conjunction with

Direct Relief and the National Association of Community Health Centers (NACHC), we announced the successful results of a four-year effort to improve access and quality of care at community health centers across the U.S., and we launched a new effort focused on enhancing the medication management process.

We also remain focused on supporting priority health needs that are aligned with the U.N. Sustainable Development Goals (SDGs). The BD 2020 Sustainability Goals are one of many ways we contribute to the SDGs through our focus on collaborating with the public and non-profit sectors, developing innovative products, expanding access to quality health services, driving environmental stewardship and empowering our associates. We are making solid progress toward our 2020 Sustainability Goals and look forward to sharing more highlights in the coming months.

The health of our planet impacts the health of humans, so we continue to take action and form partnerships to tackle environmental challenges, such as reducing greenhouse gas emissions across our value chain. We are taking measures to increase climate resilience

and investing in renewable energy, and I'm proud to report that 98 percent of electric power used by our operations in the U.S. is obtained from renewable sources.

Fiscal year 2017 financial results

Fiscal 2017 once again proved the strength of your investment in BD as we continued to deliver consistent results in line with our commitments. We grew revenues after accounting for divestitures, drove underlying margin expansion and delivered strong adjusted earnings per share (EPS) growth—all while overcoming multiple headwinds.

In highlighting some of our key achievements in fiscal 2017, I would first point to the successful transformation of the U.S. dispensing business model. Our strategy for end-to-end medication management is resonating with customers, and we are continuing to see sales growth as customers understand the benefits of our integrated offering.

Second, emerging markets continue to be a key growth driver for BD. Our momentum in China and the broader emerging markets is reflected in our strong fiscal 2017 performance in both segments.

Third, we are extremely pleased with the excellent progress we continue to make with our CareFusion cost synergies. We continue to expect approximately \$350 million dollars in total CareFusion cost synergies as we exit fiscal year 2018.

Finally and perhaps most exciting, we are making excellent progress in preparing for the integration of BD and Bard. We are energized by the opportunities we will have to bring more comprehensive, clinically relevant solutions to customers and patients around the globe.

Creating a new healthcare leader

Bringing together Bard and BD will create a new healthcare leader—one of the five largest medical technology companies in the world. And while the scale of the new BD will create an even more valuable partner for our customers, scale alone doesn't signify strength.

The power of the two companies stems from the combined focus of each legacy organization and bringing together different, yet very complementary approaches and cultures. Historically, BD has largely focused on process improvement in healthcare. By improving clinician, laboratory scientist and researchers' workflows, we support improved outcomes, safety and efficiency, all while reducing costs.

Bard has built its legacy by focusing on disease states and providing some of the best clinical solutions and treatments for areas such as peripheral vascular disease, urology, hernia and cancer, which complements the similar focus BD has on diabetes care. Both companies have long believed in the power of data to demonstrate how improvements in clinical outcomes reduce the cost of care.

By combining Bard's clinical focus on disease states with BD's focus on process improvement, we will build a world-class medical technology leader that becomes an invaluable resource for hospitals and health



systems. As healthcare providers continue to consolidate, they are looking for partners that not only bring products and solutions but the expertise around the delivery of care and the treatment of disease, from both a clinical and economical perspective. BD and Bard bring this unique combination.

Creating a new healthcare industry leader requires strong leaders as part of the management team. Last year, we promoted Tom Polen to President of BD, overseeing both the Medical and Life Sciences segments, Americas sales and R&D. He will also oversee the Interventional segment as Bard joins BD. Following the completion of the Bard transaction, Samarat (Sam) Khichi will serve as Executive Vice President and General Counsel, leading an expert Law Group for BD. Sam will join us from Bard to replace our dear friend and colleague Jeff Sherman, who passed away in 2017 after a courageous battle with cancer. We very much look forward to welcoming Sam and more than two dozen other senior leaders from Bard to our leadership team upon close.

The ability to continue to strengthen our leadership team is a testament to the company's focus on cultivating and developing talent across BD and our commitment to inclusion and diversity. I'm very proud to say that BD—for the first time ever—earned a perfect score in the Human Rights Campaign Foundation's 2018 Corporate Equality Index and was designated a "Best Place to Work for LGBTQ Equality."

Through all of this growth and change, one thing remains constant at BD—our unwavering commitment to doing what is right. We are guided by our core set of values that help ensure we don't put company results above company principles. This is one area that we will never change.

Fiscal 2017 was another transformative year for BD, and I am truly looking forward to what lies ahead. As much as BD has been able to affect positive change in healthcare for the past 120 years, we continue to strengthen our ability and leadership in our industry. At BD, we will continue to drive forward and innovate because we know there is a patient at the end of everything we do, counting on us. Thank you for your continued investment and belief in our purpose of *advancing the world of health*. We have much to look forward to together.

Sincerely,

Vincent A. Forlenza

Chairman and
Chief Executive Officer

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

Form 10-K

ANNUAL REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934
FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2017
COMMISSION FILE NUMBER 1-4802

BECTON, DICKINSON AND COMPANY
(Exact name of registrant as specified in its charter)

New Jersey
(State or other jurisdiction of incorporation or organization)

22-0760120
(I.R.S. Employer Identification No.)

1 Becton Drive
Franklin Lakes, New Jersey
(Address of principal executive offices)

07417-1880
(Zip code)

Registrant's telephone number, including area code **(201) 847-6800**

Securities registered pursuant to Section 12(b) of the Act:

<u>Title of Each Class</u>	<u>Name of Each Exchange on Which Registered</u>
Common Stock, par value \$1.00	New York Stock Exchange
Depository Shares, each representing a 1/20th interest in a share of 6.125% Cumulative Preferred Stock Series A	New York Stock Exchange
0.368% Notes due June 9, 2019	New York Stock Exchange
1.000% Notes due December 15, 2022	New York Stock Exchange
1.900% Notes due December 15, 2026	New York Stock Exchange

Securities registered pursuant to Section 12(g) of the Act:

None

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act. Yes No

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the Act. Yes No

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes No

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

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K is not contained herein, and will not be contained, to the best of registrant's knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K.

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, 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 Accelerated filer
Non-accelerated filer (Do not check if a smaller reporting company)
Smaller reporting company
Emerging growth company

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.

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Act). Yes No

As of March 31, 2017, the aggregate market value of the registrant's outstanding common stock held by non-affiliates of the registrant was approximately \$39,070,060,303.

As of October 31, 2017, 227,978,328 shares of the registrant's common stock were outstanding.

Documents Incorporated by Reference

Portions of the registrant's Proxy Statement for the Annual Meeting of Shareholders to be held January 23, 2018 are incorporated by reference into Part III hereof.

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PART I

Item 1. *Business.*

General

Becton, Dickinson and Company (also known as “BD”) was incorporated under the laws of the State of New Jersey in November 1906, as successor to a New York business started in 1897. BD’s executive offices are located at 1 Becton Drive, Franklin Lakes, New Jersey 07417-1880, and its telephone number is (201) 847-6800. All references in this Form 10-K to “BD” refer to Becton, Dickinson and Company and its domestic and foreign subsidiaries, unless otherwise indicated by the context.

BD is a global medical technology company engaged in the development, manufacture and sale of a broad range of medical supplies, devices, laboratory equipment and diagnostic products used by healthcare institutions, life science researchers, clinical laboratories, the pharmaceutical industry and the general public. We provide customer solutions that are focused on improving medication management and patient safety; supporting infection prevention practices; equipping surgical and interventional procedures; improving drug delivery; aiding anesthesiology care; enhancing the diagnosis of infectious diseases and cancers; advancing cellular research and applications; and supporting the management of diabetes.

Business Segments

BD’s operations consist of two worldwide business segments: BD Medical and BD Life Sciences. Information with respect to BD’s business segments is included in Note 6 to the consolidated financial statements contained in Item 8, Financial Statements and Supplementary Data, and is incorporated herein by reference.

BD Medical

BD Medical produces a broad array of medical technologies and devices that are used to help improve healthcare delivery in a wide range of settings. The primary customers served by BD Medical are hospitals and clinics; physicians’ office practices; consumers and retail pharmacies; governmental and nonprofit public health agencies; pharmaceutical companies; and healthcare workers. BD Medical consists of the following organizational units:

<u>Organizational Unit</u>	<u>Principal Product Lines</u>
Diabetes Care	Syringes, pen needles and other products related to the injection or infusion of insulin and other drugs used in the treatment of diabetes.
Medication and Procedural Solutions	Needles, syringes and intravenous catheters for medication delivery (including safety-engineered and auto-disable devices); prefilled IV flush syringes; regional anesthesia needles and trays; sharps disposal containers; closed-system transfer devices; skin antiseptic products; and surgical and laproscopic instrumentation.
Medication Management Solutions	Intravenous medication safety and infusion therapy delivery systems, including infusion pumps and dedicated disposables; medication compounding workflow systems; automated medication dispensing; automated supply management systems; medication inventory optimization and tracking systems; and analytics related to all the above products.
Pharmaceutical Systems	Prefillable drug delivery systems provided to pharmaceutical companies for use as containers for injectable pharmaceutical products, which are then placed on the market as drug/device combinations.

BD Life Sciences

BD Life Sciences provides products for the safe collection and transport of diagnostics specimens, and instruments and reagent systems to detect a broad range of infectious diseases, healthcare-associated infections (“HAIs”) and cancers. In addition, BD Life Sciences produces research and clinical tools that facilitate the study of cells, and the components of cells, to gain a better understanding of normal and disease processes. That information is used to aid the discovery and development of new drugs and vaccines, and to improve the diagnosis and management of diseases. The primary customers served by BD Life Sciences are hospitals, laboratories and clinics; blood banks; healthcare workers; public health agencies; physicians’ office practices; academic and government institutions; and pharmaceutical and biotechnology companies. BD Life Sciences consists of the following organizational units:

<u><i>Organizational Unit</i></u>	<u><i>Principal Product Lines</i></u>
Preanalytical Systems	Integrated systems for specimen collection; and safety-engineered blood collection products and systems.
Diagnostic Systems	Automated blood culturing and tuberculosis culturing systems; molecular testing systems for infectious diseases and women’s health; microorganism identification and drug susceptibility systems; liquid-based cytology systems for cervical cancer screening; rapid diagnostic assays; microbiology laboratory automation; and plated media.
Biosciences	Fluorescence-activated cell sorters and analyzers; monoclonal antibodies and kits for performing cell analysis; reagent systems for life science research; molecular indexing and next-generation sequencing sample preparation for genomics research; clinical oncology, immunological (HIV) and transplantation diagnostic/monitoring reagents and analyzers; and cell culture media supplements for biopharmaceutical manufacturing.

Acquisitions

Definitive Agreement to Acquire C. R. Bard, Inc.

On April 23, 2017, BD entered into a definitive agreement (the “Merger Agreement”) under which BD will acquire C.R. Bard, Inc. (“Bard”) to create a highly differentiated medical technology company uniquely positioned to improve both the process of care and the treatment of disease for patients and healthcare providers.

Under the terms of the Merger Agreement, each outstanding share of Bard common stock will be converted into the right to receive \$222.93 in cash, without interest, and 0.5077 of a share of BD’s common stock. The transaction is subject to regulatory approvals, as well as customary closing conditions, and is expected to close in the fourth calendar quarter of 2017. BD plans to finance the transaction with the issuance of BD’s common stock to Bard’s shareholders and available cash on hand, which will include net proceeds raised in the third quarter through equity and debt transactions.

The foregoing description of the Merger Agreement and the transactions contemplated thereby is not complete and is subject to and qualified in its entirety by reference to the Merger Agreement, a copy of which is included as an exhibit to the Current Report on Form 8-K filed by BD on April 24, 2017.

Acquisition of CareFusion Corporation

On March 17, 2015, BD completed the acquisition of CareFusion Corporation (“CareFusion”), a global medical technology company with a comprehensive portfolio of products in the areas of medication management, infection prevention, operating room and procedural effectiveness, and respiratory care. The CareFusion acquisition positioned BD as a global leader in medication management.

Acquisition of remaining interest in Caesarea Medical Electronics

Upon its acquisition of CareFusion, BD acquired a 40% ownership interest in Caesarea Medical Electronics ("CME"), an Israeli-based global infusion pump systems manufacturer. On April 3, 2017, BD acquired the remaining 60% ownership interest in CME.

Additional information regarding these acquisitions is contained in Note 9 to the consolidated financial statements contained in Item 8, Financial Statements and Supplementary Data, which is incorporated herein by reference.

Divestiture

In March 2016, BD signed a definitive agreement to sell 50.1% of its Respiratory Solutions business and form a joint venture with respect to this business. The Respiratory Solutions business was acquired in the CareFusion acquisition in 2015 and was a component of the Medical segment. Upon closing of the transaction, which occurred on October 3, 2016, the Company transferred the Respiratory Solutions business to a new standalone entity, retaining a 49.9% non-controlling interest in the new entity. The buyer controls the operations and governance of the new entity. Additional information regarding this transaction is contained in Note 10 to the consolidated financial statements contained in Item 8, Financial Statements and Supplementary Data, which is incorporated herein by reference.

International Operations

BD's products are manufactured and sold worldwide. For reporting purposes, we organize our operations outside the United States as follows: Europe, EMA (which includes the Commonwealth of Independent States, the Middle East and Africa); Greater Asia (which includes Japan and Asia Pacific); Latin America (which includes Mexico, Central America, the Caribbean and South America); and Canada. The principal products sold by BD outside the United States are hypodermic needles and syringes; insulin syringes and pen needles; BD Hypak™ brand prefilled syringe systems; infusion therapy products including Alaris™ infusion pumps; pharmacy automation equipment including Pyxis™ systems; BD Vacutainer™ brand blood collection products; diagnostic systems and laboratory equipment and products; flow cytometry instruments and reagents. BD has manufacturing operations outside the United States in Bosnia and Herzegovina, Brazil, Canada, China, Dominican Republic, France, Germany, Hungary, India, Ireland, Israel, Italy, Japan, Mexico, the Netherlands, Singapore, Spain, and the United Kingdom. Geographic information with respect to BD's operations is included under the heading "Geographic Information" in Note 6 to the consolidated financial statements included in Item 8, Financial Statements and Supplementary Data, and is incorporated herein by reference.

Foreign economic conditions and exchange rate fluctuations have caused the profitability related to foreign revenues to fluctuate more than the profitability related to domestic revenues. BD believes its activities in some countries outside the United States involve greater risk than its domestic business due to the factors cited herein, as well as the economic environment, local commercial and economic policies and political uncertainties. See further discussion of this risk in Item 1A. Risk Factors.

Distribution

BD's products are marketed and distributed in the United States and internationally through independent distribution channels, and directly to end-users by BD and independent sales representatives. Order backlog is not material to BD's business inasmuch as orders for BD products generally are received and filled on a current basis, except for items temporarily out of stock. BD's worldwide sales are not generally seasonal, with the exception of certain medical devices in the Medication and Procedural Solutions organizational unit, and diagnostic products in the Diagnostic Systems organizational unit, which relate to seasonal diseases such as influenza.

Raw Materials and Components

BD purchases many different types of raw materials and components, including plastics, glass, metals, textiles, paper products, agricultural products, electronic and mechanical sub-assemblies and various biological, chemical and petrochemical products. BD seeks to ensure continuity of supply by securing multiple options for sourcing. However, there are situations where raw materials and components are only available from one supplier, which are referred to as sole sourced. The use of sole sourced materials and components may be due to sourcing of proprietary and/or patented technology and processes that are intended to provide a unique market differentiation to our product. In other cases, while a raw material or component can be sourced from multiple manufacturers, only one supplier is qualified due to quality assurance, cost or other considerations. In order to provide alternate sources, BD must complete a rigorous qualification process, which most often includes completion of regulatory registration and approval. If clinical trials are not required, this qualification process can take 3-18 months depending on the criticality of the change. When clinical trials are required, this process may lengthen the qualification phase from one to three years. BD continuously assesses its sole sourced raw materials and components, and maintains business continuity plans with our suppliers. BD's continuity plans may include securing secondary supply with alternate suppliers, qualification of alternate manufacturing facilities, maintaining contingency stock, internal development of supply and establishment of technology escrow accounts. While BD works closely with its suppliers, no assurance can be given that these efforts will be successful, and there may be events that cause supply interruption, reduction or termination that adversely impacts BD's ability to manufacture and sell certain products.

Research and Development

BD conducts its research and development ("R&D") activities at its operating units and at BD Technologies in Research Triangle Park, North Carolina. The majority of BD's R&D activities are conducted in North America. Outside North America, BD primarily conducts R&D activities in China, France, India, Ireland and Singapore. BD also collaborates with certain universities, medical centers and other entities on R&D programs and retains individual consultants and partners to support its efforts in specialized fields. BD spent approximately \$774 million, \$828 million and \$632 million on research and development during the fiscal years ended September 30, 2017, 2016, and 2015, respectively.

Intellectual Property and Licenses

BD owns significant intellectual property, including patents, patent applications, technology, trade secrets, know-how, copyrights and trademarks in the United States and other countries. BD is also licensed under domestic and foreign patents, patent applications, technology, trade secrets, know-how, copyrights and trademarks owned by others. In the aggregate, these intellectual property assets and licenses are of material importance to BD's business. BD believes, however, that no single patent, technology, trademark, intellectual property asset or license is material in relation to BD's business as a whole, or to any business segment.

Competition

BD operates in the increasingly complex and challenging medical technology marketplace. Technological advances and scientific discoveries have accelerated the pace of change in medical technology, the regulatory environment of medical products is becoming more complex and vigorous, and economic conditions have resulted in a challenging market. Companies of varying sizes compete in the global medical technology field. Some are more specialized than BD with respect to particular markets, and some have greater financial resources than BD. New companies have entered the field, particularly in the areas of molecular diagnostics, safety-engineered devices and in the life sciences, and established companies have diversified their business activities into the medical technology area. Other firms engaged in the distribution of medical technology products have become manufacturers of medical devices and instruments as well. Acquisitions and collaborations by and among companies seeking a competitive advantage also affect the competitive environment. In addition, the entry into the market of low-cost manufacturers are creating increased pricing pressures. Some competitors have also established manufacturing sites or have contracted with suppliers located in these countries as a means to lower their costs.

BD competes in this evolving marketplace on the basis of many factors, including price, quality, innovation, service, reputation, distribution and promotion. The impact of these factors on BD's competitive position varies among BD's various product offerings. In order to remain competitive in the industries in which it operates, BD continues to make investments in research and development, quality management, quality improvement, product innovation and productivity improvement in support of its core strategies.

Third-Party Reimbursement

A majority of BD's customers rely on third-party payers, including government programs and private health insurance plans, to reimburse some or all of the cost of the procedures, products and services they provide. Our technologies are subject to worldwide regulations regarding reimbursement developed by government agencies, including the Centers for Medicare and Medicaid Services (CMS) in the United States; the National Health Service in the United Kingdom; the Joint Federal Committee in Germany; the Commission d'Evaluation des Produits et prestations in France; the Ministry for Health, Labor and Welfare in Japan; the Ministry of Health and the National Development and Reform Commission in China; among many others. In addition, our technologies are also subject to reimbursement policies issued by private insurance companies and managed care organizations.

BD is actively engaged in identifying and communicating value propositions of its products for payer, provider, and patient stakeholders, and it employs various efforts and resources to attempt to positively impact coverage, coding and payment pathways. However, BD has no direct control over payer decision-making with respect to coverage and payment levels for BD products. The manner and level of reimbursement in any given case may depend on the site of care, the procedure(s) performed, the final patient diagnosis, the device(s) and/or drug(s) utilized, the available budget, or a combination of these factors, and coverage and payment levels are determined at each payer's discretion. As BD's product offerings are diverse across a variety of healthcare settings, they are affected to varying degrees by the many payment pathways that impact the decisions of healthcare providers regarding which medical products they purchase and the prices they are willing to pay for those products. Therefore, changes in reimbursement levels or methods may either positively or negatively impact sales of BD products in any given country for any given product.

As government programs seek to expand healthcare coverage for their citizens, they have at the same time sought to control costs by limiting the amount of reimbursement they will pay for particular procedures, products or services. Many third-party payers have developed specific payment and delivery mechanisms to support these cost control efforts and to focus on paying for value. These mechanisms include payment reductions, pay for performance measures, quality-based performance payments, restrictive coverage policies, bidding and tender mechanics, studies to compare the effectiveness of therapies and use of technology assessments. These changes have created an increased emphasis on the delivery of more cost-effective and quality-driven healthcare.

In addition, as a result of the Patient Protection and Affordable Care Act ("PPACA"), the U.S. is implementing value based payment methodologies and seeking to create alternative payment models such as bundled payments to continue to drive improved value. We see other governments around the world considering similar bundling reform measures, including the development of the Diagnosis Related Group ("DRG") as a payment mechanism to drive toward quality and resource based reimbursement.

Regulation

BD's medical technology products and operations are subject to regulation by the U.S. Food and Drug Administration ("FDA") and various other federal and state agencies, as well as by foreign governmental agencies. These agencies enforce laws and regulations that govern the development, testing, manufacturing, labeling, advertising, marketing and distribution, and market surveillance of BD's medical products. The scope of the activities of these agencies, particularly in the Europe, Japan, and Asia Pacific regions in which BD operates, has been increasing.

BD actively maintains FDA/ISO Quality Systems that establish standards for its product design, manufacturing, and distribution processes. Prior to marketing or selling most of its products, BD must secure approval from the FDA and counterpart non-U.S. regulatory agencies. Following the introduction of a product, these agencies engage in periodic reviews and inspections of BD's quality systems, as well as product performance and advertising and promotional materials. These regulatory controls, as well as any changes in FDA policies, can affect the time and cost associated with the development, introduction and continued availability of new products. Where possible, BD anticipates these factors in its product development and planning processes. These agencies possess the authority to take various administrative and legal actions against BD, such as product recalls, product seizures and other civil and criminal sanctions. BD also undertakes voluntary compliance actions, such as voluntary recalls.

BD also is subject to various federal and state laws, and laws outside the United States, concerning healthcare fraud and abuse (including false claims laws and anti-kickback laws), global anti-corruption, transportation, safety and health, and customs and exports. Many of the agencies enforcing these laws have increased their enforcement activities with respect to medical device manufacturers in recent years. This appears to be part of a general trend toward increased regulation and enforcement activity within and outside the United States.

In addition, as part of PPACA, the federal government has enacted the Sunshine Act provisions requiring BD to publicly report gifts and payments made to physicians and teaching hospitals. Failure to comply with these provisions could result in a range of fines, penalties and/or other sanctions.

Our infusion pump organizational unit is operating under an amended consent decree entered into by CareFusion with the FDA in 2007. CareFusion's consent decree with the FDA related to its Alaris™ SE infusion pumps. In February 2009, CareFusion and the FDA amended the consent decree to include all infusion pumps manufactured by or for CareFusion 303, Inc., the organizational unit that manufactures and sells infusion pumps in the United States. The amended consent decree does not apply to intravenous administration sets and accessories.

While this BD organizational unit remains subject to the amended consent decree, which includes the requirements of the original consent decree, it has made substantial progress in its compliance efforts. However, we cannot predict the outcome of this matter, and the amended consent decree authorizes the FDA, in the event of any violations in the future, to order us to cease manufacturing and distributing infusion pumps, recall products and take other actions. We may be required to pay damages of \$15,000 per day per violation if we fail to comply with any provision of the amended consent decree, up to \$15 million per year.

We also cannot currently predict whether additional monetary investment will be incurred to resolve this matter or the matter's ultimate impact on our business. We may be obligated to pay more costs in the future because, among other things, the FDA may determine that we are not fully compliant with the amended consent decree and therefore impose penalties under the amended consent decree, and/or we may be subject to future proceedings and litigation relating to the matters addressed in the amended consent decree. As of September 30, 2017, we do not believe that a loss is probable in connection with the amended consent decree, and accordingly, we have no accruals associated with compliance with the amended consent decree.

For further discussion of risks relating to the regulations to which we are subject, see Item 1A. Risk Factors.

Employees

As of September 30, 2017, BD had 41,933 employees, of which 17,497 were employed in the U.S. (including Puerto Rico). BD believes that its employee relations are satisfactory.

Available Information

BD maintains a website at www.bd.com. BD also makes available its Annual Reports on Form 10-K, its Quarterly Reports on Form 10-Q, and its Current Reports on Form 8-K (and amendments to those reports) as soon as reasonably practicable after those reports are electronically filed with, or furnished to, the Securities and Exchange Commission (“SEC”). These filings may be obtained and printed free of charge at www.bd.com/investors. In addition, the written charters of the Audit Committee, the Compensation and Management Development Committee, the Corporate Governance and Nominating Committee, the Executive Committee and the Science, Marketing, Innovation and Technology Committee of the Board of Directors, BD’s Corporate Governance Principles and its Code of Conduct, are available and may be printed free of charge at BD’s website at www.bd.com/investors/corporate_governance/. Printed copies of these materials, this 2017 Annual Report on Form 10-K, and BD’s reports and statements filed with, or furnished to, the SEC, may also be obtained, without charge, by contacting the Corporate Secretary, BD, 1 Becton Drive, Franklin Lakes, New Jersey 07417-1880, telephone 201-847-6800. In addition, the SEC maintains an internet site that contains reports, proxy and information statements, and other information regarding issuers that file electronically with the SEC at www.sec.gov.

BD also routinely posts important information for investors on its website at www.bd.com/investors. BD may use this website as a means of disclosing material, non-public information and for complying with its disclosure obligations under Regulation FD adopted by the SEC. Accordingly, investors should monitor the Investor Relations portion of BD’s website noted above, in addition to following BD’s press releases, SEC filings, and public conference calls and webcasts. Our website and the information contained therein or connected thereto shall not be deemed to be incorporated into this Annual Report.

Forward-Looking Statements

BD and its representatives may from time-to-time make certain forward-looking statements in publicly-released materials, both written and oral, including statements contained in filings with the SEC and in its reports to shareholders. Additional information regarding BD’s forward-looking statements is contained in Item 7, Management’s Discussion and Analysis of Financial Condition and Results of Operations.

Item 1A. Risk Factors.

An investment in BD involves a variety of risks and uncertainties. The following describes some of the significant risks that could adversely affect BD’s business, financial condition, operating results or cash flows.

Risks Relating to BD

A downturn in global economic conditions could adversely affect our operations.

Deterioration in the global economic environment, particularly in emerging markets and countries with government-sponsored healthcare systems, may cause decreased demand for our products and services and increased competition, which could result in lower sales volume and downward pressure on the prices for our products, longer sales cycles, and slower adoption of new technologies. A weakening of macroeconomic conditions may also adversely affect our suppliers, which could result in interruptions in supply. We have also previously experienced delays in collecting government receivables in certain countries in Western Europe due to economic conditions, and we may experience similar delays in the future in these and other countries or regions experiencing financial problems.

The medical technology industry is very competitive.

We are a global company that faces significant competition from a wide range of companies. These include large medical device companies with multiple product lines, some of which may have greater financial and marketing resources than we do, and firms that are more specialized than we are with respect to particular markets or product lines. We face competition across all our product lines and in each market in which our products are sold on the basis of product features, clinical or economic outcomes, product quality, price, services and other factors. In addition, we face changing customer preferences and requirements, including increased customer demand for more environmentally-friendly products.

The medical technology industry is also subject to rapid technological change and discovery. The development of new or improved products, processes or technologies by other companies (such as needle-free injection technology) may render our products or proposed products obsolete or less competitive. In some instances, competitors, including pharmaceutical companies, also offer, or are attempting to develop, alternative therapies for disease states that may be delivered without a medical device. The entry into the market of manufacturers located in China and other low-cost manufacturing locations has also created pricing pressure, particularly in developing markets.

We are subject to foreign currency exchange risk.

A substantial amount of our revenues are derived from international operations, and we anticipate that a significant portion of our sales will continue to come from outside the U.S. in the future. The revenues we report with respect to our operations outside the United States may be adversely affected by fluctuations in foreign currency exchange rates. A discussion of the financial impact of exchange rate fluctuations and the ways and extent to which we may attempt to address any impact is contained in Item 7., Management's Discussion of Financial Condition and Results of Operations. Any hedging activities we engage in may only offset a portion of the adverse financial impact resulting from unfavorable changes in foreign currency exchange rates. We cannot predict with any certainty changes in foreign currency exchange rates or the degree to which we can mitigate these risks.

Changes in reimbursement practices of third-party payers could affect the demand for our products and the prices at which they are sold.

Our sales depend, in part, on the extent to which healthcare providers and facilities are reimbursed by government authorities, private insurers and other third-party payers for the costs of our products. The coverage policies and reimbursement levels of third-party payers, which can vary among public and private sources and by country, may affect which products customers purchase and the prices they are willing to pay for those products in a particular jurisdiction. Reimbursement rates can also affect the acceptance rate of new technologies and products. Legislative or administrative reforms to reimbursement systems in the United States or abroad, changes in reimbursement rates by private payers, or adverse decisions relating to our products by administrators of these systems could significantly reduce reimbursement for procedures using our products or result in denial of reimbursement for those products, which would adversely affect customer demand or the price customers are willing to pay for such products. See "Third-Party Reimbursement" under Item 1., Business.

The reinstatement of the PPACA's medical device tax may adversely affect our results of operations.

The PPACA imposes on medical device manufacturers, such as BD, a 2.3% excise tax on U.S. sales of certain medical devices. While the excise tax has been suspended until the end of 2017, absent further legislative action, it will be reinstated in 2018.

Consolidation in the healthcare industry could adversely affect our future revenues and operating income.

The medical technology industry has experienced a significant amount of consolidation, resulting in companies with greater market presence. Health care systems and other health care companies are also consolidating, resulting in greater purchasing power for these companies. As a result, competition among medical device suppliers to provide goods and services has increased. Group purchasing organizations and integrated health delivery networks have also served to concentrate purchasing decisions for some customers, which has led to downward pricing pressure for medical device suppliers. Further consolidation in the industry could intensify competition among medical device suppliers and exert additional pressure on the prices of our products.

Cost volatility could adversely affect our operations.

Our results of operations could be negatively impacted by volatility in the cost of raw materials, components, freight and energy that increases the costs of producing and distributing our products. New laws or regulations adopted in response to climate change could also increase energy costs as well as the costs of certain raw materials and components. In particular, we purchase supplies of resins, which are oil-based components used in the manufacture of certain products, and any significant increases in resin costs could adversely impact future operating results. Increases in oil prices can also increase our packaging and transportation costs. We may not be able to offset any increases in these operational costs.

Breaches of our information technology systems could have a material adverse effect on our operations.

We rely on information technology systems to process, transmit, and store electronic information in our day-to-day operations, including sensitive personal information and proprietary or confidential information. In addition, some of our products include information technology that collects data regarding patients and patient therapy on behalf of our customers and some connect to our systems for maintenance purposes. Our information technology systems have been subjected to attack via malicious code execution, cyber- or phishing-attacks, and we have experienced instances of unauthorized access to our systems in the past and expect to be subject to similar attacks in the future. In addition to our own information, in the course of doing business, we sometimes store information with third parties that could be subject to these types of attacks. Cyber-attacks could result in our intellectual property and other confidential information being accessed or stolen. Likewise, we could suffer disruption of our operations and other significant negative consequences including increased costs for security measures or remediation, diversion of management attention, and adverse impact on our relationships with vendors, business partners and customers. Unauthorized tampering, adulteration or interference with our products may also create issues with product functionality that could result in a loss of data, risk to patient safety, and product recalls or field actions. Cyber-attacks could result in unauthorized access to our systems and products which could also result in actions by regulatory bodies or civil litigation. While we will continue to dedicate significant resources to protect the company against unauthorized access to our systems and work with government authorities to detect and reduce the risk of future cyber incidents, cyber-attacks are becoming more sophisticated frequent, and adaptive. There can be no assurances that our protective measures will prevent future attacks that could have a material adverse impact on our business.

Our future growth is dependent in part upon the development of new products, and there can be no assurance that such products will be developed.

A significant element of our strategy is to increase revenue growth by focusing on innovation and new product development. New product development requires significant investment in research and development, clinical trials and regulatory approvals. The results of our product development efforts may be affected by a number of factors, including our ability to anticipate customer needs, innovate and develop new products and technologies, successfully complete clinical trials, obtain regulatory approvals and reimbursement in the United States and abroad, manufacture products in a cost-effective manner, obtain appropriate intellectual property protection for our products, and gain and maintain market acceptance of our products. In addition, patents attained by others can preclude or delay our commercialization of a product. There can be no assurance that any products now in development or that we may seek to develop in the future will achieve technological feasibility, obtain regulatory approval or gain market acceptance.

We cannot guarantee that any of our strategic acquisitions, investments or alliances will be successful.

We may seek to supplement our internal growth through strategic acquisitions, investments and alliances. Such transactions are inherently risky, and the integration of any newly-acquired business requires significant effort and management attention. The success of any acquisition, investment or alliance may be affected by a number of factors, including our ability to properly assess and value the potential business opportunity or to successfully integrate any business we may acquire into our existing business. There can be no assurance that any past or future transaction will be successful.

The international operations of our business may subject us to certain business risks.

A substantial amount of our sales come from our operations outside the United States, and we intend to continue to pursue growth opportunities in foreign markets, especially in emerging markets. Our foreign operations subject us to certain risks, including the effects of fluctuations in foreign currency exchange (discussed above), the effects of local economic and political conditions, U.S. relations with the governments of the foreign countries in which we operate, foreign regulatory requirements or changes in such requirements, local product preferences and product requirements, longer payment terms for account receivables than we experience in the U.S., difficulty in establishing, staffing and managing foreign operations, differing labor regulations, potential changes to international trade agreements and treaties, changes in tax laws, weakening or loss of the protection of intellectual property rights in some countries, import or export licensing requirements, trade protection measures and restrictions on the transfer of capital across borders. The success of our operations outside the United States depends, in part, on our ability to acquire or form and maintain alliances with local companies and make necessary infrastructure enhancements to, among other things, our production facilities and sales and distribution networks.

In addition, our international operations are governed by the U.S. Foreign Corrupt Practices Act and similar anti-corruption laws outside the U.S. Global enforcement of anti-corruption laws has increased substantially in recent years, with more enforcement proceedings by U.S. and foreign governmental agencies and the imposition of significant fines and penalties. While we have implemented policies and procedures to enhance compliance with these laws, our international operations, which often involve customer relationships with foreign governments, create the risk that there may be unauthorized payments or offers of payments made by employees, consultants, sales agents or distributors. Any alleged or actual violations of these laws may subject us to government investigations, significant criminal or civil sanctions and other liabilities, and negatively affect our reputation.

Under the U.S. tax code, we may also be subject to additional taxation to the extent we repatriate earnings from our foreign operations to the U.S. In the event we require more capital in the United States than is generated by our U.S. operations to fund acquisitions or other activities and elect to repatriate earnings from foreign jurisdictions, our effective tax rate may be higher as a result. Recently in the United States, there have been legislative proposals to tax profits that are earned abroad. These, and other proposals for fundamental U.S. corporate tax reform, if enacted, could have a material impact on our financial results.

The June 2016 referendum result in the United Kingdom (“UK”) to exit the European Union (“EU”) (commonly known as “Brexit”) and the subsequent triggering by the UK government in March 2017 of Article 50 of the Lisbon Treaty, which commenced the UK’s official withdrawal process from the EU, has created uncertainties affecting business operations in the UK and the EU. Following the referendum, there was a significant decline in the value of the British pound compared to the U.S. dollar, and continued volatility in exchange rates and economic conditions is expected as the UK negotiates its exit from the EU. Until the terms of the UK’s exit from the EU are determined, it is difficult to predict its impact but it is possible that the withdrawal could, among other things, affect the legal and regulatory schemes to which our businesses are subject, impact trade between the UK and the EU and other parties and create economic and political uncertainty in the region.

There have also been recent proposals for the U.S. to significantly modify or withdraw from certain existing international trade agreements, including the North American Free Trade Agreement. While we cannot predict whether any such changes will be implemented, it is possible that changes to international trade agreements could materially impact our business.

Reductions in customers' research budgets or government funding may adversely affect our business.

We sell products to researchers at pharmaceutical and biotechnology companies, academic institutions, government laboratories and private foundations. Research and development spending of our customers can fluctuate based on spending priorities and general economic conditions. A number of these customers are also dependent for their funding upon grants from U.S. government agencies, such as the U.S. National Institutes of Health ("NIH") and agencies in other countries. The level of government funding of research and development is unpredictable. For instance, there have been instances where NIH grants have been frozen or otherwise unavailable for extended periods. The availability of governmental research funding may be adversely affected by economic conditions and governmental spending reductions. Any reduction or delay in governmental funding could cause our customers to delay or forego purchases of our products.

A reduction or interruption in the supply of certain raw materials and components would adversely affect our manufacturing operations and related product sales.

We purchase many different types of raw materials and components used in our products. Certain raw materials and components are not available from multiple sources. In addition, for quality assurance, cost-effectiveness and other reasons, we elect to purchase certain raw materials and components from sole suppliers. The supply of these materials can be disrupted for a number of reasons, including economic conditions as described above. While we work with suppliers to ensure continuity of supply, no assurance can be given that these efforts will be successful. In addition, due to regulatory requirements relating to the qualification of suppliers, we may not be able to establish additional or replacement sources on a timely basis or without excessive cost. The termination, reduction or interruption in supply of these sole-sourced raw materials and components could adversely impact our ability to manufacture and sell certain of our products.

Interruption of our manufacturing operations could adversely affect our future revenues and operating income.

We have manufacturing sites all over the world. In some instances, however, the manufacturing of certain of our product lines is concentrated in one or more of our plants. Damage to one or more of these facilities from weather or natural disasters, or issues in our manufacturing process, equipment failure or other factors, could adversely affect our ability to manufacture these products, resulting in lost revenues and damage to our relationships with customers. In particular, damage to our manufacturing facilities in Puerto Rico resulting from Hurricane Maria in September 2017 could adversely impact our revenue and earnings results for fiscal year 2018.

We are subject to lawsuits.

We are or have been a defendant in a number of lawsuits, including purported class action lawsuits for alleged antitrust violations and suits alleging patent infringement, and could be subject to additional lawsuits in the future. A more detailed description of these lawsuits is contained in note 5 to the consolidated financial statements included in Item 8., Financial Statements and Supplementary Data. Given the uncertain nature of litigation generally, we are not able in all cases to estimate the amount or range of loss that could result from an unfavorable outcome of the litigation to which we are a party. In view of these uncertainties, we could incur charges in excess of any currently established accruals and, to the extent available, excess liability insurance. Any such future charges, individually or in the aggregate, could have a material adverse effect on our results of operations and cash flows.

We are subject to extensive regulation.

Our operations are global and are affected by complex state, federal and international laws relating to healthcare, environmental protection, antitrust, anti-corruption, marketing, fraud and abuse (including anti-kickback and false claims laws), export control, employment and other areas. Violations of these laws can result in criminal or civil sanctions, including substantial fines and, in some cases, exclusion from participation in health care programs such as Medicare and Medicaid. Environmental laws, particularly with respect to the emission of greenhouse gases, are also becoming more stringent throughout the world, which may increase our costs of operations or necessitate changes in our manufacturing plants or processes or those of our suppliers, or result in liability to BD. The enactment of additional laws in the future may increase our compliance costs or otherwise adversely impact our operations.

We are also subject to extensive regulation by the FDA pursuant to the Federal Food, Drug and Cosmetic Act, by comparable agencies in foreign countries, and by other regulatory agencies and governing bodies. Most of our products must receive clearance or approval from the FDA or counterpart regulatory agencies in other countries before they can be marketed or sold. The process for obtaining marketing approval or clearance may take a significant period of time and require the expenditure of substantial resources, and these have been increasing due to increased requirements from the FDA for supporting data for submissions. The process may also require changes to our products or result in limitations on the indicated uses of the products. Governmental agencies may also impose new requirements regarding registration, labeling or prohibited materials that may require us to modify or re-register products already on the market or otherwise impact our ability to market our products in those countries. Once clearance or approval has been obtained for a product, there is an obligation to ensure that all applicable FDA and other regulatory requirements continue to be met. Following the introduction of a product, these agencies also periodically review our manufacturing processes and product performance. Our failure to comply with the applicable good manufacturing practices, adverse event reporting, clinical trial and other requirements of these agencies could delay or prevent the production, marketing or sale of our products and result in fines, delays or suspensions of regulatory clearances, closure of manufacturing sites, seizures or recalls of products and damage to our reputation. More stringent oversight by the FDA and other agencies in recent years has resulted in increased enforcement activity, which increases the compliance risk for us and other companies in our industry.

As a result of the CareFusion acquisition, we are operating under a consent decree with the FDA that was entered into by CareFusion in 2009, that affects our infusion pump business in the United States. For more information regarding the consent decree, see “Regulation” under Item 1, “Business”.

Defects or quality issues associated with our products could adversely affect the results of our operations.

The design, manufacture and marketing of medical devices involve certain inherent risks. Manufacturing or design defects, unapproved use of our products, or inadequate disclosure of risks or other information relating to the use of our products can lead to injury or other adverse events. These events could lead to recalls or safety alerts relating to our products (either voluntary or required by the FDA or similar governmental authorities in other countries), and could result, in certain cases, in the removal of a product from the market. A recall could result in significant costs and lost sales and customers, enforcement actions and/or investigations by state and federal governments or other enforcement bodies, as well as negative publicity and damage to our reputation that could reduce future demand for our products. Personal injuries relating to the use of our products can also result in significant product liability claims being brought against us. In some circumstances, such adverse events could also cause delays in regulatory approval of new products.

Our operations are dependent in part on patents and other intellectual property assets.

Many of our businesses rely on patent, trademark and other intellectual property assets. These intellectual property assets, in the aggregate, are of material importance to our business. We can lose the protection afforded by these intellectual property assets through patent expirations, legal challenges or governmental action. Patents attained by competitors, particularly as patents on our products expire, may also adversely affect our competitive position. In addition, competitors may seek to invalidate patents on our products or claim that our products infringe upon their intellectual property, which could result in a loss of competitive advantage or the payment of significant legal fees, damage awards and past or future royalties, as well as injunctions against future sales of our products. We also operate in countries that do not protect intellectual property rights to the same extent as in the U.S., which could make it easier for competitors to compete with us in those countries. The loss of a significant portion of our portfolio of intellectual property assets may have an adverse effect on our earnings, financial condition or cash flows.

Natural disasters, war and other events could adversely affect our future revenues and operating income.

Natural disasters (including pandemics), war, terrorism, labor disruptions and international conflicts, and actions taken by the United States and other governments or by our customers or suppliers in response to such events, could cause significant economic disruption and political and social instability in the United States and areas outside of the United States in which we operate. These events could result in decreased demand for our products, adversely affect our manufacturing and distribution capabilities, or increase the costs for or cause interruptions in the supply of materials from our suppliers.

We need to attract and retain key employees to be competitive.

Our ability to compete effectively depends upon our ability to attract and retain executives and other key employees, including people in technical, marketing, sales and research positions. Competition for experienced employees, particularly for persons with specialized skills, can be intense. Our ability to recruit such talent will depend on a number of factors, including compensation and benefits, work location and work environment. If we cannot effectively recruit and retain qualified executives and employees, our business could be adversely affected.

We may not realize all of the anticipated benefits and cost savings resulting from our acquisition of CareFusion.

While we have realized significant cost savings to date in connection with our acquisition of CareFusion in 2015, achieving additional cost synergies may prove more difficult than expected, and it is possible that the anticipated cost synergies of the merger may not be realized fully, or may take longer to realize than expected.

Risks Relating To Our Acquisition of Bard

Completion of the Bard acquisition is subject to conditions and if these conditions are not satisfied or waived, the Bard acquisition will not be completed.

The obligations of us and Bard to complete the Bard acquisition are subject to satisfaction or waiver of a number of conditions, the expiration or termination of the applicable waiting period in connection with the Hart-Scott-Rodino Antitrust Improvements Act of 1976, as amended (the “HSR Act”), the receipt of any authorization or consent from certain other governmental authorities required to be obtained with respect to the merger under applicable foreign antitrust laws, approval of the listing on the NYSE of shares of our common stock to be issued in the Bard acquisition, and the absence of an injunction prohibiting the Bard acquisition. Each party’s obligation to complete the Bard acquisition is subject to the satisfaction or waiver (to the extent permitted under applicable law) of certain other customary conditions, the accuracy of the representations and warranties of the other party under the Merger Agreement (subject to the materiality standards set forth in the Merger Agreement), the performance by the other party of its respective obligations under the Merger Agreement in all material respects and delivery of officer certificates by the other party certifying satisfaction of the two preceding conditions. Either we or Bard may, subject to certain exceptions, terminate the Merger Agreement upon mutual consent or if the Bard acquisition has not been consummated on or before January 23, 2018 (or before April 23, 2018 if all closing conditions have been satisfied other than the receipt of required competition approvals).

The failure to satisfy all of the required conditions could delay the completion of the Bard acquisition for a significant period of time or prevent it from occurring. If the Bard acquisition is not completed, our ongoing business may be materially adversely affected and, without realizing any of the benefits of having completed the Bard acquisition, we will be subject to a number of risks, including the following:

- the market price of our common stock could decline;
- if the Merger Agreement is terminated and our board of directors seeks another business combination, our stockholders cannot be certain that we will be able to find a party willing to enter into a transaction on terms equivalent to or more attractive than the terms that Bard has agreed to in the Merger Agreement;
- time and resources, financial and other, committed by our management to matters relating to the Bard acquisition could otherwise have been devoted to pursuing other beneficial opportunities for our company;

- we may experience negative reactions from the financial markets or from our customers or employees; and
- we will be required to pay our respective costs relating to the Bard acquisition, including legal, accounting, financial advisory, financing and printing fees, whether or not the Bard acquisition is completed.

In addition, if the Bard acquisition is not completed, we could be subject to litigation related to any failure to complete the Bard acquisition or related to any enforcement proceeding commenced against us to perform our obligations under the Merger Agreement. The materialization of any of these risks could materially and adversely impact our ongoing business.

Similarly, any delay in completing the Bard acquisition could, among other things, result in additional transaction costs, loss of revenue or other negative effects associated with uncertainty about completion of the Bard acquisition and cause us not to realize some or all of the benefits that we expect to achieve if the Bard acquisition is successfully completed within its expected timeframe. There can be no assurance that the conditions to the closing of the Bard acquisition will be satisfied or waived or that the Bard acquisition will be consummated.

In order to complete the Bard acquisition, we and Bard must make certain governmental filings and obtain certain governmental authorizations, and if such filings and authorizations are not made or granted or are granted with conditions, completion of the Bard acquisition may be jeopardized or the anticipated benefits of the Bard acquisition could be reduced.

Although we and Bard have agreed in the Merger Agreement to use reasonable best efforts, subject to certain limitations, to make certain governmental filings, to obtain the required expiration or termination of the waiting period under the HSR Act and to obtain any authorization or consent from certain other governmental authorities required to be obtained with respect to the merger under applicable foreign antitrust laws, there can be no assurance that such approvals will be obtained. As a condition to granting termination of the waiting period under the HSR Act and to adoption of approvals of the Bard acquisition, governmental authorities may impose requirements, limitations or costs or require divestitures or place restrictions on the conduct of our business after completion of the Bard acquisition.

Under the terms of the Merger Agreement, subject to certain exceptions, we and our subsidiaries are required to accept certain conditions and take certain actions imposed by certain governmental authorities that would apply to, or affect, the businesses, assets or properties of us, our subsidiaries or Bard and its subsidiaries. There can be no assurance that regulators will not impose conditions, terms, obligations or restrictions and that such conditions, terms, obligations or restrictions will not have the effect of (i) delaying completion of the Bard acquisition, (ii) imposing additional material costs on or materially limiting the revenues of the combined company following the Bard acquisition, or (iii) otherwise adversely affecting our businesses and results of operations after completion of the Bard acquisition. In addition, we can provide no assurance that these conditions, terms, obligations or restrictions will not result in the delay or abandonment of the Bard acquisition.

Each party is subject to business uncertainties and contractual restrictions while the proposed merger is pending, which could adversely affect each party's or the combined company's business and operations.

In connection with the pendency of the Bard acquisition, it is possible that some customers, suppliers and other persons with whom we or Bard have a business relationship may delay or defer certain business decisions or might decide to seek to terminate, change or renegotiate their relationships with us or Bard, as the case may be, as a result of the Bard acquisition, which could negatively affect our or Bard's respective revenues, earnings and cash flows, regardless of whether the Bard acquisition is completed. If the Bard acquisition is completed, such terminations, changes or renegotiations could negatively affect the revenues, earnings and cash flows of the combined company. These risks may be exacerbated by delays or other adverse developments with respect to the completion of the Bard acquisition.

Combining the two companies may be more difficult, costly or time consuming than expected and the anticipated benefits and cost savings of the Bard acquisition may not be realized.

We and Bard have operated and, until the completion of the Bard acquisition, will continue to operate, independently. The success of the Bard acquisition, including anticipated benefits and cost savings, will depend, in part, on our ability to successfully combine and integrate our business with the business of Bard.

The Bard acquisition will involve the integration of Bard's business with our existing business, which is a complex, costly and time-consuming process. It is possible that the pendency of the Bard acquisition and/or the integration process could result in material challenges, including, without limitation:

- the diversion of management's attention from ongoing business concerns and performance shortfalls at one or both of the companies as a result of the devotion of management's attention to the Bard acquisition;
- managing a larger combined company;
- maintaining employee morale and retaining key management and other employees;
- the possibility of faulty assumptions underlying expectations regarding the integration process;
- retaining existing business and operational relationships and attracting new business and operational relationships;
- consolidating corporate and administrative infrastructures and eliminating duplicative operations and inconsistencies in standards, controls, procedures and policies;
- coordinating geographically separate organizations;
- unanticipated issues in integrating information technology, communications and other systems; and
- unforeseen expenses or delays associated with the Bard acquisition.

Many of these factors will be outside of the combined company's control and any one of them could result in delays, increased costs, decreases in revenues and diversion of management's time and energy, which could materially affect the combined company's financial position, results of operations and cash flows.

If we experience difficulties with the integration process, the anticipated benefits of the Bard acquisition may not be realized fully or at all, or may take longer to realize than expected. These integration matters could have an adverse effect on (i) each of us and Bard during this transition period and (ii) the combined company for an undetermined period after completion of the Bard acquisition. In addition, the actual cost savings of the Bard acquisition could be less than anticipated.

In addition, certain risks associated with our industry and business described herein and in our public filings may become more significant following consummation of the Bard acquisition, including, but not limited to, risks relating to: the continued focus by third-party payors on cost containment and government scrutiny of the healthcare industry's sales and marketing practices, various healthcare reform proposals that have emerged on the federal and state levels and in other jurisdictions where the combined company sells its products, collective bargaining and labor activity, and the integrity of our information systems that are run by third party vendors and such vendors' ability to maintain their systems and reduce any vulnerability to natural and system disruptions and prevent cyber-attacks and other unauthorized access.

The future results of the combined company may be adversely impacted if the combined company does not effectively manage its expanded operations following the completion of the Bard acquisition.

Following the completion of the Bard acquisition, the size of the combined company's business will be significantly larger than the current size of either our or Bard's respective businesses. The combined company's ability to successfully manage this expanded business will depend, in part, upon management's ability to design and implement strategic initiatives that address not only the integration of two discrete companies, but also the increased scale and scope of the combined business with its associated increased costs and complexity. There can be no assurances that the combined company will be successful or that it will realize the expected operating efficiencies, cost savings and other benefits currently anticipated from the Bard acquisition.

The combined company is expected to incur substantial expenses related to the completion of the Bard acquisition and the integration of BD and Bard.

We and Bard have incurred, and expect to continue to incur, a number of non-recurring costs associated with the Bard acquisition and combining the operations of the two companies. The substantial majority of non-recurring expenses will be comprised of transaction and regulatory costs related to the Bard acquisition.

We also will incur transaction fees and costs related to formulating and implementing integration plans, including facilities and systems consolidation costs and employment-related costs. We continue to assess the magnitude of these costs, and additional unanticipated costs may be incurred in the Bard acquisition and the integration of the two companies' businesses. Although we expect that the elimination of duplicative costs, as well as the realization of other efficiencies related to the integration of the businesses, should allow us to offset integration-related costs over time, this net benefit may not be achieved in the near term, or at all.

In connection with the Bard acquisition, we have incurred significant additional indebtedness, and certain of Bard's indebtedness will remain outstanding, which could adversely affect us, including by decreasing our business flexibility, and will increase our interest expense.

We have substantially increased our indebtedness in connection with the pending Bard acquisition through the incurrence of new indebtedness to finance the Bard acquisition and, following the Bard acquisition, through the assumption of Bard's existing indebtedness, in comparison to our indebtedness on a recent historical basis, which could have the effect of, among other things, reducing our flexibility to respond to changing business and economic conditions and increasing our interest expense.

The amount of cash required to pay interest on our increased indebtedness levels following completion of the Bard acquisition, and thus the demands on our cash resources, will be greater than the amount of cash flows required to service our indebtedness prior to the Bard acquisition. The increased levels of indebtedness following completion of the Bard acquisition could also reduce funds available for working capital, capital expenditures, acquisitions, the repayment or refinancing of our indebtedness as it becomes due and other general corporate purposes and may create competitive disadvantages for us relative to other companies with lower debt levels. In addition, certain of the indebtedness incurred in connection with the Bard acquisition bears interest at variable interest rates. If interest rates increase, variable rate debt will create higher debt service requirements, which could further adversely affect our cash flows. If we do not achieve the expected benefits and cost savings from the Bard acquisition, or if the financial performance of the combined company does not meet current expectations, then our ability to service our indebtedness may be adversely impacted.

In addition, our credit ratings affect the cost and availability of future borrowings and, accordingly, our cost of capital. Our ratings reflect each rating organization's opinion of our financial strength, operating performance and ability to meet our debt obligations. There can be no assurance that we will achieve a particular rating or maintain a particular rating in the future or that we will be able to maintain our current rating. Furthermore, we expect that our combined company's credit ratings will be lower following the Bard acquisition, including below "investment grade" by Moody's Investors Service, Inc., which may further increase the combined company's future borrowing costs and reduce the combined company's access to capital.

Moreover, in the future we may be required to raise substantial additional financing to fund working capital, capital expenditures, the repayment or refinancing of our indebtedness, acquisitions or other general corporate requirements. Our ability to arrange additional financing or refinancing will depend on, among other factors, our financial position and performance, as well as prevailing market conditions and other factors beyond our control. We cannot assure you that it will be able to obtain additional financing or refinancing on terms acceptable to us or at all.

We may not be able to service all of the combined company's indebtedness and may be forced to take other actions to satisfy our obligations under our indebtedness, which may not be successful. Our failure to meet our debt service obligations could have a material adverse effect on our business, financial condition and results of operations.

We depend on cash on hand and cash flows from operations to make scheduled debt payments. We expect to be able to meet the estimated cash interest payments on the combined company's debt following the Bard acquisition through a combination of the expected cash flows from operations of the combined company. However, our ability to generate sufficient cash flow from operations of the combined company and to utilize other methods to make scheduled payments will depend on a range of economic, competitive and business factors, many of which are outside of our control. There can be no assurance that these sources will be adequate. If we are unable to service our indebtedness and fund our operations, we will be forced to reduce or delay capital expenditures, seek additional capital, sell assets or refinance our indebtedness. Any such action may not be successful and we may be unable to service our indebtedness and fund our operations, which could have a material adverse effect on our business, financial condition or results of operations.

The agreements that will govern the indebtedness incurred in connection with the Bard acquisition contain various covenants that impose restrictions on us and certain of our subsidiaries that may affect our ability to operate our businesses.

The agreements that govern the indebtedness incurred in connection with the Bard acquisition contain various affirmative and negative covenants that may, subject to certain significant exceptions, restrict the ability of certain of our subsidiaries to incur debt and the ability of us and certain of our subsidiaries to, among other things, have liens on our property, and/or merge or consolidate with any other person or sell or convey certain of our assets to any one person, engage in certain transactions with affiliates and change the nature of our business. In addition, the agreements also require us to comply with certain financial covenants, including financial ratios. Our ability and the ability of our subsidiaries to comply with these provisions may be affected by events beyond our control. Failure to comply with these covenants could result in an event of default, which, if not cured or waived, could accelerate our repayment obligations and could result in a default and acceleration under other agreements containing cross-default provisions. Under these circumstances, we might not have sufficient funds or other resources to satisfy all of our obligations.

Uncertainties associated with the Bard acquisition may cause a loss of management personnel and other key employees of Bard or us, which could adversely affect the future business and operations of the combined company following the Bard acquisition.

We and Bard are dependent on the experience and industry knowledge of our respective officers and other key employees to execute our respective business plans. The combined company's success after the Bard acquisition will depend in part upon its ability to retain key management personnel and other key employees of us and Bard. Current and prospective employees of us and Bard may experience uncertainty about their future roles with the combined company following the Bard acquisition, which may materially adversely affect the ability of each of us and Bard to attract and retain key personnel during the pendency of and after the Bard acquisition. Accordingly, no assurance can be given that the combined company will be able to retain key management personnel and other key employees of us and Bard.

Completion of the Bard acquisition will trigger change in control or other provisions in certain agreements to which Bard is a party, which may have an adverse impact on the combined company's business and results of operations.

The completion of the Bard acquisition will trigger change in control and other provisions in certain agreements to which Bard is a party. If we and Bard are unable to negotiate waivers of those provisions, the counterparties may exercise their rights and remedies under the agreements, potentially terminating the agreements or seeking monetary damages. Even if we and Bard are able to negotiate waivers, the counterparties may require a fee for such waivers or seek to renegotiate the agreements on terms less favorable to Bard or the combined company. Any of the foregoing or similar developments may have an adverse impact on the combined company's business and results of operations.

For example, if the ratings of certain of Bard's outstanding senior notes are reduced beyond certain thresholds within certain time periods prior to or following the consummation of the Bard acquisition, Bard could be required to offer to repurchase such notes at 101% of the aggregate principal amount of such notes plus any accrued and unpaid interest to the repurchase date.

Following the consummation of the Bard acquisition, the combined company will assume certain potential liabilities relating to Bard, including certain products liability and mass torts claims.

Following the consummation of the Bard acquisition, the combined company will have assumed certain potential liabilities relating to Bard, including certain products liability and mass tort claims with respect to the design, manufacture and marketing of medical devices and related settlement agreements and judgments. As of September 30, 2017, Bard has reported that there are: (i) approximately 25 federal and 185 state lawsuits involving individual claims by approximately 205 plaintiffs, as well as one putative class action in the United States, are currently pending against Bard's hernia repair implant products, (ii) product liability lawsuits involving individual claims by approximately 3,285 plaintiffs are currently pending against Bard in various federal and state jurisdictions with respect to Bard's surgical continence products for women and (iii) product liability lawsuits involving individual claims by approximately 1,755 plaintiffs are currently pending against Bard in various federal and state jurisdictions with respect to Bard's vena cava filter products.

Bard does not maintain or has limited remaining insurance coverage for certain of these claims and the combined company may not be able to obtain additional insurance on acceptable terms or at all that will provide adequate protection against potential liabilities. Moreover, in some circumstances adverse events arising from or associated with the design, manufacture, quality or marketing of our combined company's products could result in the FDA suspending or delaying its review of our applications for new product approvals, or imposing post market approval requirements. In addition, reserves established by Bard or the combined company for estimated losses, including with respect to these claims, do not represent an exact calculation of actual liability but instead represent estimates of the probable loss at the time the reserve is established. Due to the inherent uncertainty underlying loss reserve estimates, additional reserves may be established from time-to-time, and actual losses relating to the assumed Bard liabilities may be materially higher or lower than the related reserve. Any of the foregoing could have a material adverse effect on our business, financial condition or results of operations.

Sales of shares of BD common stock after the completion of the transaction may cause the market price of BD equity securities to fall.

We will issue a significant number of shares of our common stock in connection with the Bard acquisition. Many Bard stockholders may decide not to hold the shares of our common stock they will receive in the Bard acquisition. Other Bard stockholders, such as funds with limitations on their permitted holdings of stock in individual issuers, may be required to sell the shares of our common stock that they receive in the Bard acquisition. Such sales of our common stock could have the effect of depressing the market price for our equity securities and may take place promptly following the Bard acquisition.

The mandatory convertible preferred stock underlying the depositary shares issued in connection with the financing of the Bard transaction may adversely affect the market price of BD common stock.

The market price of BD common stock is likely to be influenced by the mandatory convertible preferred stock underlying the depositary shares issued in connection with the financing for the Bard transaction. The market price of BD common stock could become more volatile and could be depressed by:

- investors' anticipation of the potential resale in the market of a substantial number of additional shares of BD common stock received upon conversion of the mandatory convertible preferred stock;
- possible sales of BD common stock by investors who view the mandatory convertible preferred stock as a more attractive means of equity participation in BD than owning shares of BD common stock; and
- hedging or arbitrage trading activity that may develop involving the mandatory convertible preferred stock and BD common stock.

Item 1B. *Unresolved Staff Comments.*

None.

Item 2. Properties.

BD's executive offices are located in Franklin Lakes, New Jersey. As of October 31, 2017, BD owned or leased 289 facilities throughout the world, comprising approximately 20,462,405 square feet of manufacturing, warehousing, administrative and research facilities. The U.S. facilities, including those in Puerto Rico, comprise approximately 7,472,419 square feet of owned and 2,976,267 square feet of leased space. The international facilities comprise approximately 7,478,714 square feet of owned and 2,535,005 square feet of leased space. Sales offices and distribution centers included in the total square footage are also located throughout the world.

Operations in each of BD's business segments are conducted at both U.S. and international locations. Particularly in the international marketplace, facilities often serve more than one business segment and are used for multiple purposes, such as administrative/sales, manufacturing and/or warehousing/distribution. BD generally seeks to own its manufacturing facilities, although some are leased. The following table summarizes property information by business segment.

Sites	Corporate	BD Life Sciences	BD Medical	Mixed(A)	Total
Leased	14	25	96	83	218
Owned	6	26	33	6	71
Total	20	51	129	89	289
Square feet	2,263,694	4,421,732	10,838,632	2,938,347	20,462,405

(A) Facilities used by more than one business segment.

BD believes that its facilities are of good construction and in good physical condition, are suitable and adequate for the operations conducted at those facilities, and are, with minor exceptions, fully utilized and operating at normal capacity.

The U.S. facilities are located in Alabama, Arizona, California, Connecticut, Florida, Georgia, Illinois, Indiana, Maryland, Massachusetts, Michigan, Missouri, Nebraska, New Jersey, North Carolina, Ohio, Oklahoma, South Carolina, Texas, Utah, Virginia, Washington, D.C., Washington, Wisconsin and Puerto Rico.

The international facilities are as follows:

- *Europe, Middle East, Africa*, which includes facilities in Austria, Belgium, Bosnia and Herzegovina, the Czech Republic, Denmark, England, Finland, France, Germany, Ghana, Hungary, Ireland, Israel, Italy, Kenya, Luxembourg, Netherlands, Norway, Poland, Portugal, Russia, Saudi Arabia, South Africa, Spain, Sweden, Switzerland, Turkey, the United Arab Emirates and Zambia.

- *Greater Asia*, which includes facilities in Australia, Bangladesh, China, India, Indonesia, Japan, Malaysia, New Zealand, the Philippines, Singapore, South Korea, Taiwan, Thailand and Vietnam.

- *Latin America*, which includes facilities in Argentina, Brazil, Chile, Colombia, Mexico, Peru and the Dominican Republic.

- *Canada*.

Item 3. Legal Proceedings.

Information with respect to certain legal proceedings is included in Note 5 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data, and is incorporated herein by reference.

Item 4. Mine Safety Disclosures.

Not applicable.

Executive Officers of the Registrant

The following is a list of the executive officers of BD, their ages and all positions and offices held by each of them during the past five years. There is no family relationship between any executive officer or director of BD.

Name	Age	Position
Vincent A. Forlenza	64	Chairman since July 2012; Chief Executive Officer since October 2011; President from January 2009 to April 2017; and Chief Operating Officer from July 2010 to October 2011.
Thomas E. Polen	44	President since April 2017; Executive Vice President and President - Medical Segment from October 2014 to April 2017; Group President from October 2013 to October 2014; and Worldwide President - BD Diagnostic Systems from October 2010 to October 2013.
James W. Borzi	55	Executive Vice President, Global Operations and Chief Supply Chain Office since October 2017; Senior Vice President, Global Operations from 2015 to October 2017; Vice President, Global Manufacturing from 2013 to 2015; and Vice President and General Manager, Hydro Aluminum from 2012 to 2013.
Alexandre Conroy	54	Worldwide President, Medication and Procedural Solutions since May 2017; and Executive Vice President and President, Europe, EMA and the Americas from June 2012 to May 2017.
Roland Goette	55	Executive Vice President and President, EMEA since May 2017; President, Europe from October 2014 to May 2017; and prior thereto, Vice President and General Manager - Medical Surgical Systems, Western Europe.
James Lim	53	Executive Vice President and President, Greater Asia since June 2012.
Alberto Mas	56	Executive Vice President and President - Life Sciences Segment since October 2016; Worldwide President - Life Sciences, Diagnostic Systems from October 2013 to October 2016; and Worldwide President - BD Biosciences from October 2011 to October 2013.
Christopher R. Reidy	60	Executive Vice President, Chief Financial Officer and Chief Administrative Officer since July 2013; and prior thereto, Vice President and Chief Financial Officer of ADP Corporation.
Nabil Shabshab	52	Worldwide President, Diabetes Care and Digital Health since August 2017; Executive Vice President and President, Americas and Chief Customer Experience Officer from May 2017 to August 2017; Executive Vice President and Chief Marketing Officer from January 2015 to May 2017; Senior Vice President and Chief Marketing Officer from August 2011 to January 2015.
Ellen R. Strahlman, M.D.	60	Executive Vice President, Research and Development since January 2015, Chief Medical Officer since April 2013; Senior Vice President, Research and Development from April 2013 to January 2015; and prior thereto, Senior Vice President, Office of the CEO and Global Head, Neglected Tropical Diseases of GlaxoSmithKline.
Linda M. Tharby	49	Executive Vice President and Chief Human Resource Officer since October 2016; Executive Vice President and President - Life Sciences Segment from October 2014 to October 2016; Group President from October 2013 to October 2014; and prior thereto, Worldwide President - BD Medical, Diabetes Care.

PART II

Item 5. *Market for Registrant's Common Equity, Related Stockholder Matters and Issuer Purchases of Equity Securities.*

BD's common stock is listed on the New York Stock Exchange. As of October 31, 2017, there were approximately 13,134 shareholders of record.

Market and Market Prices of Common Stock (per common share)

By Quarter	2016		2017	
	High	Low	High	Low
First	\$156.53	\$132.19	\$179.17	\$162.80
Second	152.54	132.88	185.34	164.80
Third	172.19	152.86	195.15	177.07
Fourth	181.55	169.64	205.63	191.56

Dividends (per common share)

By Quarter	2016	2017
First	\$ 0.660	\$ 0.730
Second	0.660	0.730
Third	0.660	0.730
Fourth	0.660	0.730

Issuer Purchases of Equity Securities

The table below sets forth certain information regarding BD's purchases of its common stock during the fiscal quarter ended September 30, 2017.

Period	Total Number of Shares Purchased(1)	Average Price Paid per Share	Total Number of Shares Purchased as Part of Publicly Announced Plans or Programs(2)	Maximum Number of Shares that May Yet be Purchased Under the Plans or Programs(2)
July 1-31, 2017	1,809	\$197.71	—	7,857,742
August 1-31, 2017	240	\$196.39	—	7,857,742
September 1-30, 2017	—	—	—	7,857,742
Total	2,049	\$197.55	—	7,857,742

- (1) Includes 2,049 shares purchased during the quarter in open market transactions by the trust relating to BD's Deferred Compensation and Retirement Benefit Restoration Plan and 1996 Directors' Deferral Plan.
- (2) Represents shares available under the repurchase program authorized by the Board of Directors on September 24, 2013 for 10 million shares, for which there is no expiration date.

Item 6. Selected Financial Data.

FIVE-YEAR SUMMARY OF SELECTED FINANCIAL DATA
Becton, Dickinson and Company

	Years Ended September 30				
	2017	2016	2015	2014	2013
	Dollars in millions, except share and per share amounts				
Operations					
Revenues	\$ 12,093	\$ 12,483	\$ 10,282	\$ 8,446	\$ 8,054
Gross Margin	5,942	5,991	4,695	4,301	4,171
Research and Development Expense	774	828	632	550	494
Operating Income	1,478	1,430	1,074	1,606	1,254
Interest Expense, Net	445	367	356	89	98
Income From Continuing Operations Before Income Taxes	976	1,074	739	1,522	1,165
Income Tax (Benefit) Provision	(124)	97	44	337	236
Income from Continuing Operations	1,100	976	695	1,185	929
Net Income	1,100	976	695	1,185	1,293
Basic Earnings Per Share from Continuing Operations	4.70	4.59	3.43	6.13	4.76
Diluted Earnings Per Share from Continuing Operations	4.60	4.49	3.35	5.99	4.67
Dividends Per Common Share	2.92	2.64	2.40	2.18	1.98
Financial Position					
Total Current Assets	\$ 18,633	\$ 6,367	\$ 5,659	\$ 5,775	\$ 5,530
Total Current Liabilities	3,342	4,400	4,381	2,225	2,122
Total PPE, Net	4,638	3,901	4,060	3,605	3,476
Total Assets	37,734	25,586	26,478	12,384	12,029
Total Long-Term Debt	18,667	10,550	11,370	3,768	3,763
Total Shareholders' Equity	12,948	7,633	7,164	5,053	5,043
Book Value Per Common Share	56.80	35.79	34.00	26.33	25.99
Financial Relationships					
Gross Profit Margin	49.1%	48.0%	45.7%	50.9%	51.8%
Return on Revenues	9.1%	7.8%	6.8%	14.0%	11.5% (A)
Return on Total Assets(B)	4.7%	5.6%	5.7%	13.6%	11.1% (A)
Return on Equity	10.7%	13.2%	11.4%	23.5%	20.2% (A)
Debt to Capitalization(C)	57.5%	57.2%	59.4%	43.6%	43.6% (A)
Additional Data					
Number of Employees	41,900	50,900	49,500	30,600	30,000
Number of Shareholders	13,183	13,788	14,547	8,210	8,412
Average Common and Common Equivalent Shares Outstanding — Assuming Dilution (millions)	223.6	217.5	207.5	197.7	199.2
Depreciation and Amortization	\$ 1,088	\$ 1,114	\$ 891	\$ 562	\$ 546
Capital Expenditures	727	693	596	592	522

- (A) Excludes discontinued operations.
- (B) Earnings before interest expense and taxes as a percent of average total assets.
- (C) Total debt as a percent of the sum of total debt, shareholders' equity and non-current deferred income tax liabilities.

The results above include the impact of the specified items detailed below. Additional discussion regarding the specified items in fiscal years 2017, 2016 and 2015 are provided in Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations.

Millions of dollars, except per share amounts	Years Ended September 30				
	2017	2016	2015	2014	2013
Total specified items	\$ 1,466	\$ 1,261	\$ 1,186	\$ 153	\$ 442
After-tax impact of specified items	\$ 971	\$ 892	\$ 786	\$ 101	\$ 279
Impact of specified items on diluted earnings per share	\$ (4.34)	\$ (4.10)	\$ (3.79)	\$ (0.51)	\$ (1.40)
Impact of dilution from share issuances	\$ (0.54)	\$ —	\$ (0.02)	\$ —	\$ —

Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations

The following commentary should be read in conjunction with the consolidated financial statements and accompanying notes. Within the tables presented throughout this discussion, certain columns may not add due to the use of rounded numbers for disclosure purposes. Percentages and earnings per share amounts presented are calculated from the underlying amounts. References to years throughout this discussion relate to our fiscal years, which end on September 30.

Company Overview

Description of the Company and Business Segments

Becton, Dickinson and Company ("BD") is a global medical technology company engaged in the development, manufacture and sale of a broad range of medical supplies, devices, laboratory equipment and diagnostic products used by healthcare institutions, life science researchers, clinical laboratories, the pharmaceutical industry and the general public. The Company's organizational structure is based upon two principal business segments, BD Medical ("Medical") and BD Life Sciences ("Life Sciences").

BD's products are manufactured and sold worldwide. Our products are marketed in the United States and internationally through independent distribution channels and directly to end-users by BD and independent sales representatives. We organize our operations outside the United States as follows: Europe; EMA (which includes the Commonwealth of Independent States, the Middle East and Africa); Greater Asia (which includes Japan and Asia Pacific); Latin America (which includes Mexico, Central America, the Caribbean, and South America); and Canada. We continue to pursue growth opportunities in emerging markets, which include the following geographic regions: Eastern Europe, the Middle East, Africa, Latin America and certain countries within Asia Pacific. We are primarily focused on certain countries whose healthcare systems are expanding, in particular, China and India.

Strategic Objectives

BD remains focused on delivering sustainable growth and shareholder value, while making appropriate investments for the future. BD management operates the business consistent with the following core strategies:

- To increase revenue growth by focusing on our core products, services and solutions that deliver greater benefits to patients, healthcare workers and researchers;

- To continue investment in research and development for platform extensions and innovative new products;
- To make investments in growing our operations in emerging markets;
- To improve operating effectiveness and balance sheet productivity;
- To drive an efficient capital structure and strong shareholder returns.

Our strategy focuses on four specific areas within healthcare and life sciences:

- Enabling safer, simpler and more effective parenteral drug delivery;
- Improving clinical outcomes through new, more accurate and faster diagnostics;
- Providing tools and technologies to the research community that facilitate the understanding of the cell, cellular diagnostics and cell therapy;
- Enhancing disease management in diabetes, women's health and cancer, infectious disease and other targeted conditions.

We continue to strive to improve the efficiency of our capital structure and follow these guiding principles:

- To operate the Company consistent with an investment grade credit profile;
- To ensure access to the debt market for strategic opportunities;
- To optimize the cost of capital based on market conditions.

In assessing the outcomes of these strategies as well as BD's financial condition and operating performance, management generally reviews quarterly forecast data, monthly actual results, segment sales and other similar information. We also consider trends related to certain key financial data, including gross profit margin, selling and administrative expense, investment in research and development, return on invested capital, and cash flows.

Definitive Agreement to Acquire C.R. Bard, Inc.

On April 23, 2017, we announced that we entered into a definitive agreement under which BD will acquire C. R. Bard, Inc. ("Bard") for an implied value of \$317.00 per Bard common share in cash and stock, for estimated total consideration of approximately \$24 billion. The combination will create a highly differentiated medical technology company uniquely positioned to improve both the process of care and the treatment of disease for patients and healthcare providers. Additional discussion regarding the acquisition agreement and the related financing the Company has secured, through equity and debt issuances, is provided in Notes 3, 9 and 15 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. The transaction is subject to regulatory approval and customary closing conditions, and is expected to close in the fourth calendar quarter of 2017.

Acquisition of CareFusion

On March 17, 2015, BD acquired a 100% interest in CareFusion Corporation ("CareFusion"). CareFusion's operating results were included in BD's consolidated results of operations beginning on April 1, 2015 and as such, the consolidated results of operations for the first six months of fiscal year 2015 referenced in the commentary provided further below did not include CareFusion's results. CareFusion operates as part of our Medical segment.

Summary of Financial Results

Worldwide revenues in 2017 of \$12.093 billion decreased 3.1% from the prior-year period. The decrease reflected an approximate 7% reduction in revenues due to the divestiture of the Respiratory Solutions business in October 2016. Volume growth in 2017 of more than 4% for our continuing businesses was partially offset by an unfavorable impact of foreign currency translation of less than 1%. Pricing did not materially impact 2017 revenues. Additional disclosures regarding our divestiture of the Respiratory Solutions

business are provided in Note 10 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. Volume growth in 2017 reflected the following:

- Medical segment volume growth in 2017 was driven by sales growth in all of the segment's units, particularly by growth in the Medication and Procedural Solutions, Medication Management Solutions and Pharmaceutical Systems units.
- Life Sciences segment volume growth in 2017 was driven by growth in all three of its organizational units, particularly in its Preanalytical and Diagnostic Systems units.
- U.S. volume growth in 2017 primarily reflected growth in sales in the Medical segment's Medication Management Solutions and Diabetes Care units, as well as in all of the Life Sciences segment's units.
- International volume growth in 2017 was driven by sales in the Medical segment's Medication and Procedural Solutions, Medication Management Solutions and Pharmaceutical Systems units, as well as by sales in the Life Sciences segment's Preanalytical Systems and Diagnostic Systems units.

We continue to invest in research and development, geographic expansion, and new product promotions to drive further revenue and profit growth. Our ability to sustain our long-term growth will depend on a number of factors, including our ability to expand our core business (including geographical expansion), develop innovative new products, and continue to improve operating efficiency and organizational effectiveness. While the economic environment for the healthcare industry and healthcare utilization in the United States have generally stabilized, destabilization in the future could adversely impact our businesses. Additionally, macroeconomic challenges in Europe continue to constrain healthcare utilization, although we currently view the environment as stable. In emerging markets, the Company's growth is dependent primarily on government funding for healthcare systems. In addition, pricing pressure exists for certain geographies and could adversely impact our businesses.

Our financial position remains strong, with cash flows from operating activities totaling \$2.550 billion in 2017. At September 30, 2017, we had \$14.2 billion in cash and equivalents and short-term investments, which included net proceeds raised through registered public offerings of equity securities and debt transactions during the third quarter of approximately \$4.8 billion and \$9.6 billion, respectively. We continued to return value to our shareholders in the form of dividends. During fiscal year 2017, we paid cash dividends of \$677 million. We also repurchased approximately \$220 million of our common stock during 2017.

Each reporting period, we face currency exposure that arises from translating the results of our worldwide operations to the U.S. dollar at exchange rates that fluctuate from the beginning of such period. The ongoing relative strength of the U.S. dollar resulted in an unfavorable foreign currency translation impact to our revenue and earnings growth during fiscal year 2017. We evaluate our results of operations on both a reported and a foreign currency-neutral basis, which excludes the impact of fluctuations in foreign currency exchange rates. As exchange rates are an important factor in understanding period-to-period comparisons, we believe the presentation of results on a foreign currency-neutral basis in addition to reported results helps improve investors' ability to understand our operating results and evaluate our performance in comparison to prior periods. Foreign currency-neutral ("FXN") information compares results between periods as if exchange rates had remained constant period-over-period. We use results on a foreign currency-neutral basis as one measure to evaluate our performance. We calculate foreign currency-neutral percentages by converting our current-period local currency financial results using the prior-period foreign currency exchange rates and comparing these adjusted amounts to our current-period results. These results should be considered in addition to, not as a substitute for, results reported in accordance with U.S. generally accepted accounting principles ("GAAP"). Results on a foreign currency-neutral basis, as we present them, may not be comparable to similarly titled measures used by other companies and are not measures of performance presented in accordance with U.S. GAAP.

Results of Operations

Medical Segment

The following is a summary of Medical revenues by organizational unit:

(Millions of dollars)				2017 vs. 2016			2016 vs. 2015		
	2017	2016	2015	Total Change	Estimated FX Impact	FXN Change	Total Change	Estimated FX Impact	FXN Change
Medication and Procedural Solutions	\$3,497	\$3,413	\$2,850	2.5 %	(0.7)%	3.2 %	19.8%	(3.6)%	23.4%
Medication Management Solutions (A)	2,295	2,197	1,015	4.4 %	(0.5)%	4.9 %	116.6%	(2.3)%	118.9%
Diabetes Care	1,056	1,023	1,012	3.3 %	(0.3)%	3.6 %	1.1%	(3.3)%	4.4%
Pharmaceutical Systems	1,256	1,199	1,167	4.8 %	(0.5)%	5.3 %	2.7%	(2.4)%	5.1%
Respiratory Solutions (A)	—	822	417	NM	— %	NM	97.2%	(2.3)%	99.5%
Total Medical revenues	<u>\$8,105</u>	<u>\$8,654</u>	<u>\$6,460</u>	<u>(6.4)%</u>	<u>(0.6)%</u>	<u>(5.8)%</u>	<u>34.0%</u>	<u>(3.0)%</u>	<u>37.0%</u>
Medical segment safety-engineered products	<u>\$1,960</u>	<u>\$1,924</u>	<u>\$1,499</u>	<u>1.9 %</u>	<u>(0.3)%</u>	<u>2.2 %</u>	<u>28.3%</u>	<u>(2.9)%</u>	<u>31.2%</u>

- (A) The presentation of prior-period amounts has been revised to conform with the presentation of current-period amounts, which does not separately present an immaterial adjustment for the amortization of a deferred revenue balance write-down relating to the CareFusion acquisition.
- (B) "NM" denotes that the percentage is not meaningful.

Medical segment revenue growth in 2017 was driven by the Medication and Procedural Solutions unit's sales of infusion disposables products, particularly in international markets, and the Pharmaceutical Systems unit's sales of self-injection systems. Revenue growth in 2017 also reflected the Diabetes Care unit's increased sales of pen needles in the United States and emerging markets. International growth in the Diabetes Care unit was impacted by weaker revenues in Europe, primarily in the United Kingdom, due to increasing pressure from government payers as part of austerity measures. Medical segment revenues in 2017 were unfavorably impacted by the divestiture of the Respiratory Solutions business and the modification to dispensing equipment lease contracts with customers in the Medication Management Solutions unit, which took place in April 2017. As a result of the lease modification, substantially all new lease contracts entered into beginning in April 2017 will be accounted for as operating leases with revenue recognized over the agreement term, rather than upon the placement of capital. In 2017, revenues in the Medication Management Solutions unit included \$151 million of revenues relating to amended preexisting lease contracts.

The Medical segment's growth in 2016 largely reflected the inclusion of CareFusion's sales for a full fiscal year in 2016 compared with half the fiscal year in 2015, as previously discussed. Medical segment revenue growth in 2016 additionally reflected the Medication and Procedural Solutions unit's international sales of safety-engineered products, the Diabetes Care unit's sales of pen needles and the Pharmaceutical Systems unit's sales of self-injection systems. Fiscal year 2016 Medical segment revenue growth was unfavorably impacted by the termination of a distribution contract in the Respiratory Solutions unit.

Medical segment operating income was as follows:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Medical segment operating income	\$ 2,155	\$ 2,052	\$ 1,530
<i>Segment operating income as % of Medical revenues</i>	<i>26.6%</i>	<i>23.7%</i>	<i>23.7%</i>

The Medical segment's operating income is driven by its performance with respect to gross profit margin and operating expenses. The Medical segment's gross profit margin in 2017 was higher as compared with 2016 primarily due to the divestiture of the Respiratory Solutions business, which had products with relatively lower gross profit margins. Gross profit margin in 2017 also reflected lower manufacturing costs resulting from continuous improvement projects which enhanced the efficiency of our operations. The Medical segment's gross profit margin as a percentage of revenues was slightly higher in 2016 as compared with 2015 primarily due to lower manufacturing costs resulting from continuous operations improvement projects, partially offset by the recognition of a full year of amortization relating to intangible assets acquired in the CareFusion transaction and by unfavorable foreign currency translation.

Selling and administrative expense as a percentage of revenues in 2017 was lower compared with 2016, primarily due to the divestiture of the Respiratory Solutions business, as this business generally had a lower operating margin. Selling and administrative expense as a percentage of revenues in 2016 was favorably impacted by the suspension of the medical device excise tax imposed under the U.S. Patient Protection Affordable Care Act. Research and development expense as a percentage of revenues in 2017 reflected ongoing investment in new products and platforms, but was lower compared with 2016 as expense in 2016 included a one-time payment relating to one of the segment's ongoing projects. Research and development expense as a percentage of revenues in 2016 increased from 2015, which reflected increased investment in new products and platforms, including the one-time payment noted above.

Life Sciences Segment

The following is a summary of Life Sciences revenues by organizational unit:

<u>(Millions of dollars)</u>				<u>2017 vs. 2016</u>			<u>2016 vs. 2015</u>		
	<u>2017</u>	<u>2016</u>	<u>2015</u>	<u>Total Change</u>	<u>Estimated FX Impact</u>	<u>FXN Change</u>	<u>Total Change</u>	<u>Estimated FX Impact</u>	<u>FXN Change</u>
Preanalytical Systems	\$ 1,471	\$ 1,409	\$ 1,391	4.4%	(0.8)%	5.2%	1.3 %	(3.9)%	5.2%
Diagnostic Systems	1,378	1,301	1,299	5.9%	(0.5)%	6.4%	0.1 %	(3.2)%	3.3%
Biosciences	1,139	1,119	1,132	1.8%	(0.6)%	2.4%	(1.2)%	(2.7)%	1.5%
Total Life Sciences revenues	<u>\$ 3,988</u>	<u>\$ 3,829</u>	<u>\$ 3,822</u>	<u>4.2%</u>	<u>(0.6)%</u>	<u>4.8%</u>	<u>0.2 %</u>	<u>(3.2)%</u>	<u>3.4%</u>
Life Sciences segment safety-engineered products	<u>\$ 1,167</u>	<u>\$ 1,113</u>	<u>\$ 1,097</u>	<u>4.9%</u>	<u>(0.8)%</u>	<u>5.7%</u>	<u>1.4 %</u>	<u>(3.7)%</u>	<u>5.1%</u>

The Life Sciences segment's 2017 revenues reflected growth in global sales of the Preanalytical Systems unit's safety-engineered products and growth in sales of the Diagnostics Systems unit's microbiology and molecular platforms, particularly in emerging markets. The segment's 2017 revenue growth was also driven by increased Biosciences unit sales, particularly in developed markets.

Fiscal year 2016 revenues in the Life Sciences segment were driven by the Preanalytical Systems unit's U.S. and international sales of safety-engineered products. Segment revenue growth in 2016 also reflected the Diagnostic Systems unit's sales of its core microbiology products, as well as placements of its *BD Kiestra*TM

platform. Fiscal year 2016 revenues in the Life Sciences segment additionally reflected the Biosciences unit's research instrument and reagent sales, primarily in the United States, which were partially offset by decreased sales of the Biosciences unit's HIV-related clinical products in Africa.

Life Sciences segment operating income was as follows:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Life Sciences segment operating income	\$ 772	\$ 793	\$ 839
<i>Segment operating income as % of Life Sciences revenues</i>	<i>19.4%</i>	<i>20.7%</i>	<i>21.9%</i>

The Life Sciences segment's operating income is driven by its performance with respect to gross profit margin and operating expenses. The Life Sciences segment's gross profit margin as a percentage of revenues was lower in fiscal year 2017 primarily due to unfavorable foreign currency translation, higher raw material costs and unfavorable product mix, partially offset by lower manufacturing costs resulting from operations improvement projects. The Life Sciences segment's gross profit margin as a percentage of revenues was lower in fiscal year 2016 compared with 2015 primarily due to unfavorable foreign currency translation, partially offset by lower manufacturing costs resulting from operations improvement projects.

Selling and administrative expense as a percentage of Life Sciences revenues in 2017 was higher compared to 2016 primarily due to slightly higher administrative costs. Selling and administrative expense as a percentage of Life Sciences revenues decreased in 2016 compared with 2015, primarily due to the suspension of the medical device excise tax. Research and development expense as a percentage of revenues in 2017 was relatively flat compared with 2016, which increased compared to 2015 due to increased investment in new products and platforms.

Geographic Revenues

BD's worldwide revenues by geography are provided below.

<u>(Millions of dollars)</u>				<u>2017 vs. 2016</u>			<u>2016 vs. 2015</u>		
	<u>2017</u>	<u>2016</u>	<u>2015</u>	<u>Total Change</u>	<u>Estimated FX Impact</u>	<u>FXN Change</u>	<u>Total Change</u>	<u>Estimated FX Impact</u>	<u>FXN Change</u>
United States	\$ 6,504	\$ 6,893	\$ 5,069	(5.6)%	—	(5.6)%	36.0%	—	36.0%
International	5,589	5,590	5,213	— %	(1.2)%	1.2 %	7.2%	(6.2)%	13.4%
Total revenues	<u>\$12,093</u>	<u>\$12,483</u>	<u>\$10,282</u>	<u>(3.1)%</u>	<u>(0.5)%</u>	<u>(2.6)%</u>	<u>21.4%</u>	<u>(3.1)%</u>	<u>24.5%</u>

U.S. revenues in 2017 were unfavorably impacted by the Medical segment's divestiture of the Respiratory Solutions business and the modification to dispensing equipment lease contracts with customers in the Medical segment's Medication Management Solutions unit, as previously discussed. These impacts to U.S. revenues in 2017 were partially offset by growth in sales in the Medical segment's Medication Management Solutions and Diabetes Care units, as well as in all of the Life Sciences segment's units.

U.S. revenue growth in 2016 primarily reflected the inclusion of CareFusion's U.S. sales for the full fiscal year. U.S. revenues also reflected growth in sales of the Medical segment's legacy products, particularly in the Medication and Procedural Solutions and Pharmaceutical Systems units. U.S. revenue growth in 2016 was also driven by sales in the Life Science's segment's Preanalytical Systems and Biosciences units.

International revenues in 2017 were driven by sales in the Medical segment's Medication and Procedural Solutions, Medication Management Solutions and Pharmaceutical Systems units, as well as by sales in the Life Sciences segment's Preanalytical Systems and Diagnostic Systems units. International revenue growth in 2017 was partially offset by the impact of the Medical segment's divestiture of the Respiratory Solutions business.

International revenue growth in 2016 primarily reflected the inclusion of CareFusion's sales for the full fiscal year, as well as growth attributable to the Medical segment's legacy products in the Medication and Procedural Solutions unit. International revenue growth in 2016 also reflected the Life Sciences segment's Preanalytical Systems unit's sales, primarily in Western Europe and Asia Pacific. The Life Sciences segment's international revenue growth in 2016 was negatively impacted by a decrease in certain sales in its Biosciences unit in Africa, as previously discussed.

Emerging market revenues were \$1.95 billion, \$1.9 billion and \$1.8 billion in 2017, 2016 and 2015, respectively. Emerging market revenues in 2016 related to divested businesses, primarily the Respiratory Solutions business, were approximately \$105 million. Unfavorable foreign currency translation impacted emerging market revenues in 2017 and 2016 by an estimated \$29 million and \$156 million, respectively. Emerging market revenue growth in 2017 was driven by sales in Greater Asia, including China, and Latin America. Emerging market revenue growth in 2016 reflected the inclusion of CareFusion's sales for the full fiscal year, as well as growth in China and Latin America, partially offset by declines in the Middle East and Africa.

Specified Items

Reflected in the financial results for 2017, 2016 and 2015 were the following specified items:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Integration costs ^(A)	\$ 237	\$ 192	\$ 95
Restructuring costs ^(A)	85	526	271
Transaction costs ^(A)	39	10	59
Financing costs ^(B)	131	—	107
Purchase accounting adjustments ^(C)	491	527	645
Lease contract modification-related charge ^(D)	748	—	—
Litigation-related items ^(E)	(337)	—	12
Losses on debt extinguishment ^(F)	73	—	—
Pension settlement charges	—	6	—
Other, net	—	—	(5)
Total specified items	<u>1,466</u>	<u>1,261</u>	<u>1,186</u>
Tax impact of specified items	495	369	400
After-tax impact of specified items	<u>\$ 971</u>	<u>\$ 892</u>	<u>\$ 786</u>

- (A) Represents integration, restructuring and transaction costs, recorded in *Acquisitions and other restructurings*, which are further discussed below.
- (B) The amount in 2017 represents financing costs incurred in connection with the agreement to acquire Bard, including bridge financing commitment fees of \$79 million, which were recorded in *Interest expense*. The amount in 2015 represents financing costs incurred in connection with the CareFusion acquisition, including bridge financing commitment fees.
- (C) Primarily represents non-cash amortization expense associated with acquisition-related identifiable intangible assets. BD's amortization expense is primarily recorded in *Cost of products sold*. The adjustment in 2015 also included a fair value step-up adjustment of \$293 million recorded relative to CareFusion's inventory on the acquisition date.
- (D) Represents a non-cash charge, which was recorded in *Other operating expense, net* resulting from a modification to our dispensing equipment lease contracts with customers, as further discussed below.
- (E) The amount in 2017 largely represents the reversal of certain reserves related to an appellate court decision recorded in *Other operating expense, net* as further discussed below.
- (F) Represents losses recognized in *Other (expense) income, net* upon our extinguishment of certain long-term senior notes in the first and third quarters, as further discussed below.

Gross Profit Margin

The comparison of gross profit margins in 2017 and 2016 and the comparison of gross profit margins in 2016 and 2015 reflected the following impacts:

	<u>2017</u>	<u>2016</u>
Gross profit margin % prior-year period	48.0 %	45.7 %
Operating performance	0.7 %	2.5 %
Impact of divestitures	0.8 %	— %
CareFusion acquisition-related asset depreciation and amortization	— %	0.6 %
Foreign currency translation	(0.4)%	(0.8)%
Gross profit margin % current-year period	<u>49.1 %</u>	<u>48.0 %</u>

The operating performance impacts in 2017 and 2016 primarily reflected lower manufacturing costs resulting from the continuous operations improvement projects discussed above. Gross profit margin in 2017 was favorably impacted by businesses divestitures, primarily the divestiture of the Respiratory Solutions business which had products with relatively lower gross profit margins. Gross profit margin in 2016 benefited from a favorable comparison to 2015, which reflected the fair value step-up adjustment recorded relative to CareFusion's inventory on the acquisition date, as previously discussed, partially offset by the recognition in 2016 of a full year of amortization relating to CareFusion's intangible assets.

Operating Expenses

Operating expenses in 2017, 2016 and 2015 were as follows:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>	<u>Increase (decrease) in basis points</u>	
				<u>2017 vs. 2016</u>	<u>2016 vs. 2015</u>
Selling and administrative expense	\$ 2,925	\$ 3,005	\$ 2,563		
<i>% of revenues</i>	24.2%	24.1%	24.9%	10	(80)
Research and development expense	\$ 774	\$ 828	\$ 632		
<i>% of revenues</i>	6.4%	6.6%	6.1%	(20)	50
Acquisitions and other restructurings	\$ 354	\$ 728	\$ 426		
Other operating expense, net	\$ 410	\$ —	\$ —		

Selling and administrative

Selling and administrative expense as a percentage of revenues in 2017 was relatively flat compared with 2016. Selling and administrative expense as a percentage of revenues in 2016 as compared with 2015 reflected synergies resulting from the CareFusion acquisition, as well as favorable foreign currency translation and a suspension of the medical device excise tax, as previously discussed, partially offset by higher selling expenses relating to product launches and higher shipping expenses.

Research and development

Research and development expense as a percentage of revenues was slightly lower in 2017, which reflected a favorable comparison to 2016, which included increased investment in high growth opportunities.

Acquisitions and other restructurings

Costs relating to acquisitions and other restructurings in 2017, 2016 and 2015 primarily represented integration and restructuring costs substantially associated with our fiscal year 2015 acquisition of CareFusion and other portfolio rationalization initiatives. Integration costs in 2017 also included costs relating to our pending acquisition of Bard. Restructuring costs in 2016 included a \$214 million charge recorded to impair capitalized internal-use software assets held for sale as a result of information technology function transformation efforts. Transaction costs were incurred in 2017 in connection with the pending acquisition of Bard and other portfolio rationalization initiatives. Transaction costs were also incurred in 2015 in connection with the CareFusion acquisition. For further disclosures regarding the costs relating to acquisitions and other restructurings, refer to Notes 7, 9, 10 and 11 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data.

Other operating (income) expense, net

Other operating expense in 2017 included the \$748 million non-cash charge resulting from the modification to our dispensing equipment lease contracts with customers. Additional disclosures regarding this lease contract modification are provided in Note 17 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. Other operating income in 2017 included a \$336 million reversal of certain reserves related to an appellate court decision which, among other things, reversed an unfavorable antitrust judgment in the Retractable Technologies, Inc. case. Additional disclosures regarding this legal matter are provided in Note 5 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data

Net Interest Expense

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Interest expense	\$ (521)	\$ (388)	\$ (371)
Interest income	76	21	15
Net interest expense	<u>\$ (445)</u>	<u>\$ (367)</u>	<u>\$ (356)</u>

The increase in interest expense in 2017 compared with 2016 primarily reflected higher levels of debt due to our issuances of senior unsecured U.S. notes during the third quarter of 2017, as well as bridge financing commitment fees of \$79 million. The increase in interest expense in 2016 reflected a full year of increased financing costs associated with the CareFusion acquisition, including interest on senior unsecured notes issued in December 2014. This increase in financing costs was partially offset by the full year impact of favorable amortization of the acquisition-date fair value step-up on CareFusion's long-term debt as well as by the prior-year impact of commitment fees incurred for a bridge loan facility. Additional disclosures regarding our financing arrangements and debt instruments are provided in Note 15 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data.

The increase in interest income in 2017 compared with 2016 primarily reflected higher levels of cash on hand, as a result of our third quarter issuances of debt and equity securities, as well as higher investment gains on assets related to our deferred compensation plans. The increase in interest income in 2016 reflected the realization of investment gains on assets related to our deferred compensation plans, compared with the realization of losses in 2015, partially offset by lower cash levels outside of the United States. The offsetting movements in the deferred compensation plan liability were recorded in *Selling and administrative expense*.

Income Taxes

The income tax rates in 2017, 2016 and 2015 were as follows:

	2017	2016	2015
Effective income tax rate - (benefit) provision	(12.7)%	9.1%	5.9%
<i>Favorable impact, in basis points, from specified items</i>	2,790	1,090	1,720

The decrease in the effective income tax rate in 2017 largely reflected the more favorable tax impact from specified items recognized in 2017 compared with 2016, as well as the tax benefits recorded upon the settlement of share-based compensation awards in 2017. The share-based compensation-related tax benefits were recognized in connection with BD's adoption of new accounting requirements relating to the income tax effects of share-based compensation awards. Additional disclosures regarding this adoption are provided in Note 2 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. The increase in the effective income tax rate from 2015 to 2016 is primarily due to the decrease in tax benefits on specified items as the tax benefits on specified items in 2015 were primarily incurred in higher tax jurisdictions. The income tax rates in fiscal years 2016 also reflected the extension of the U.S. research and development income tax credit, which was partially offset by the unfavorable impact of one-time discrete items.

Net Income and Diluted Earnings per Share

Net Income and Diluted Earnings per Share in 2017, 2016 and 2015 were as follows:

	2017	2016	2015
Net income (Millions of dollars)	\$ 1,100	\$ 976	\$ 695
Diluted Earnings per Share	\$ 4.60	\$ 4.49	\$ 3.35
Unfavorable impact-specified items	\$ (4.34)	\$ (4.10)	\$ (3.79)
Unfavorable impact-foreign currency translation	\$ (0.23)	\$ (0.64)	\$ (0.69)
Dilutive impact from share issuances	\$ (0.54)	\$ —	\$ (0.02)

The dilutive impact from share issuances in 2017 represents the impact of BD shares issued in the third quarter of fiscal year 2017, in anticipation of the pending Bard acquisition. The dilutive impact from share issuances in 2015 represents the impact from shares issued as consideration for the CareFusion acquisition, prior to the inclusion of CareFusion in consolidated results of operations.

Financial Instrument Market Risk

We selectively use financial instruments to manage market risk, primarily foreign currency exchange risk and interest rate risk relating to our ongoing business operations. The counterparties to these contracts are highly rated financial institutions. We do not enter into financial instruments for trading or speculative purposes.

Foreign Exchange Risk

BD and its subsidiaries transact business in various foreign currencies throughout Europe, Greater Asia, Canada and Latin America. We face foreign currency exposure from the effect of fluctuating exchange rates on payables and receivables relating to transactions that are denominated in currencies other than our functional currency. These payables and receivables primarily arise from intercompany transactions. We hedge substantially all such exposures, primarily through the use of forward contracts. We also face currency exposure that arises from translating the results of our worldwide operations, including sales, to the U.S. dollar at exchange rates that have fluctuated from the beginning of a reporting period. From time to time, we may

purchase forward contracts and options to hedge certain forecasted transactions that are denominated in foreign currencies in order to partially protect against a reduction in the value of future earnings resulting from adverse foreign exchange rate movements. Gains or losses on derivative instruments are largely offset by the gains or losses on the underlying hedged transactions. We did not enter into contracts to hedge cash flows against foreign currency fluctuations in fiscal year 2017 or 2016.

Derivative financial instruments are recorded on our balance sheet at fair value. For foreign currency derivatives, market risk is determined by calculating the impact on fair value of an assumed change in foreign exchange rates relative to the U.S. dollar. Fair values were estimated based upon observable inputs, specifically spot currency rates and foreign currency prices for similar assets and liabilities.

With respect to the foreign currency derivative instruments outstanding at September 30, 2017 and 2016, the impact changes in the U.S. dollar would have on pre-tax earnings was estimated as follows:

<u>(Millions of dollars)</u>	<u>Increase (decrease)</u>	
	<u>2017</u>	<u>2016</u>
10% appreciation in U.S. dollar	\$ (38)	\$ (67)
10% depreciation in U.S. dollar	\$ 38	\$ 67

These calculations do not reflect the impact of exchange gains or losses on the underlying transactions that would substantially offset the results of the derivative instruments.

Interest Rate Risk

Our primary interest rate risk relates to U.S. dollar borrowings which are partially offset by U.S. dollar cash investments. When managing interest rate exposures, we strive to achieve an appropriate balance between fixed and floating rate instruments. We may enter into interest rate swaps to help maintain this balance and manage debt and interest-bearing investments in tandem, since these items have an offsetting impact on interest rate exposure. For interest rate derivative instruments, fair values are measured based upon the present value of expected future cash flows using market-based observable inputs including credit risk and interest rate yield curves. Market risk for these instruments is determined by calculating the impact to fair value of an assumed change in interest rates across all maturities.

With respect to the interest rate derivatives outstanding at September 30, 2017 and 2016, a 10% change in interest rates would not materially impact the fair value of these derivatives. Based on our overall interest rate exposure at September 30, 2017 and 2016, a 10% change in interest rates would not have a material effect on our earnings or cash flows over a one-year period.

Liquidity and Capital Resources

The following table summarizes our consolidated statement of cash flows in 2017, 2016 and 2015:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Net cash provided by (used for)			
Operating activities	\$ 2,550	\$ 2,559	\$ 1,730
Investing activities	\$ (883)	\$ (669)	\$ (8,318)
Financing activities	\$ 10,977	\$ (1,761)	\$ 6,190

Net Cash Flows from Operating Activities

The fiscal year 2017, 2016 and 2015 changes in net cash provided by operating activities was primarily attributable to net income, as adjusted for depreciation and amortization and other non-cash items. The fiscal year 2017 change in operating assets and liabilities was a net use of cash and primarily reflected higher levels of prepaid expenses, trade receivables and inventory, partially offset by higher levels of accounts payable and accrued expenses. The fiscal year 2016 change in operating assets and liabilities was a net source of cash and

primarily reflected higher levels of accounts payable and accrued expenses as well as lower levels of inventory, prepayments and financing receivables, partially offset by higher levels of accounts receivables. Net cash provided by operating activities in 2017 also reflected an adjustment for the non-cash charge resulting from the modification to our dispensing equipment lease contracts with customers, as previously discussed, and the losses recorded upon our extinguishment of certain long-term notes, which are included within *Other, net*. The previously discussed non-cash charge recorded to impair capitalized internal-use software assets held for sale is included within *Other, net* in 2016. Net cash provided by operating activities in 2016 was reduced by changes in the pension obligation resulting primarily from a discretionary cash contribution of \$100 million.

Net Cash Flows from Investing Activities

Capital expenditures

Our investments in capital expenditures are focused on projects that enhance our cost structure and manufacturing capabilities, and support our strategy of geographic expansion with select investments in growing markets. Capital expenditures of \$727 million, \$693 million, \$596 million in 2017, 2016 and 2015, respectively, primarily related to manufacturing capacity expansions and details of spending by segment are contained in Note 6 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data.

Investments

Cash inflows from the sales of investments of \$840 million in 2015 were attributable to the maturities of time deposits in Europe, Latin America and Asia Pacific.

Acquisitions of Businesses

Cash outflows relating to acquisitions in 2017 included payments for acquisitions which were immaterial both individually and in the aggregate. Cash outflows relating to acquisitions in 2015 of \$8.414 billion were primarily attributable to the completion of the CareFusion acquisition in the second quarter of fiscal year 2015. For further discussion, refer to Note 9 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data.

Divestitures of Businesses

Cash inflows relating to business divestitures in 2017 were \$165 million. Net cash flows from investing activities in 2016 included \$158 million of proceeds from the sales of non-core assets.

Net Cash Flows from Financing Activities

Net cash from financing activities in 2017, 2016 and 2015 included the following significant cash flows:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Cash inflow (outflow)			
Increase/(decrease) in borrowings under commercial paper program	\$ (200)	\$ (500)	\$ 500
Issuances of senior unsecured U.S. notes	\$ 9,616	\$ —	\$ 6,164
Issuances of euro-denominated notes	\$ 1,846	\$ —	\$ —
Payments of debt	\$(3,980)	\$ (752)	\$ (6)
Issuances of equity securities	\$ 4,827	\$ —	\$ —
Share repurchases under accelerated share repurchase agreement	\$ (220)	\$ —	\$ —
Dividends paid	\$ (677)	\$ (562)	\$ (485)

Additional disclosures regarding the equity and debt-related financing activities detailed above are provided in Notes 3 and 15 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. No further share repurchases are currently planned, as our share repurchase program has been suspended in connection with the announced agreement to acquire Bard.

Debt-Related Activities

Certain measures relating to our total debt were as follows:

	<u>2017</u>	<u>2016</u>	<u>2015</u>
Total debt (Millions of dollars)	<u>\$ 18,870</u>	<u>\$ 11,551</u>	<u>\$ 12,822</u>
Short-term debt as a percentage of total debt	1.1%	8.7%	11.3%
Weighted average cost of total debt	3.3%	3.6%	3.3%
Total debt as a percentage of total capital (A)	57.5%	57.2%	59.4%

(A) Represents shareholders' equity, net non-current deferred income tax liabilities, and debt.

The decrease in short-term debt as a percentage of total debt at September 30, 2017 was largely driven by our issuance of \$9.675 billion of senior unsecured U.S. notes during the third quarter of fiscal year 2017. The ratio of short-term debt as a percentage of total debt at September 30, 2016 primarily reflected the reclassifications of \$800 million of notes from long-term debt to short-term debt, offset by the repayment of \$750 million of floating rate notes during the third quarter of fiscal year 2016 and a reduction of commercial paper borrowings. Additional disclosures regarding our debt instruments are provided in Note 15 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data.

Cash and Short-term Investments

At September 30, 2017, total worldwide cash and short-term investments were \$14.201 billion, of which \$1.196 billion was held in jurisdictions outside of the United States. Total cash at September 30, 2017 included net proceeds raised through public offerings of equity securities and debt transactions which occurred during the third quarter of fiscal year 2017, as previously discussed.

We regularly review the amount of cash and short-term investments held outside the United States and currently intend to use such amounts to fund our international operations and their growth initiatives. In addition, if these amounts were repatriated from foreign jurisdictions to the United States, there could be adverse tax consequences.

Credit Facilities

We have in place a \$1.5 billion syndicated credit facility which can be used for general corporate purposes. There were no borrowings outstanding under this credit facility at September 30, 2017. During the first quarter of fiscal year 2017, we extended the expiration date of this credit facility to January 2022 from the original expiration date of January 2021. We may issue up to \$100 million in letters of credit under this facility and it also includes a provision that enables BD, subject to additional commitments made by the lenders, to access up to an additional \$500 million in financing through the facility for a maximum aggregate commitment of \$2 billion. The credit facility includes a single financial covenant that requires BD to maintain an interest expense coverage ratio of not less than 4-to-1 for the most recent four consecutive fiscal quarters. We were in compliance with this covenant as of September 30, 2017. We also have informal lines of credit outside the United States.

The Company had no commercial paper borrowings outstanding as of September 30, 2017.

The developments discussed below occurred in 2017 relative to our credit facilities in connection with the announcement of the acquisition of Bard.

Term loan and revolving credit facilities

In May 2017, we entered into a three-year \$2.25 billion senior unsecured term loan facility. The proceeds from this facility may only be used to fund a portion of the cash consideration for the Bard acquisition, as well as the fees and expenses incurred in connection with this acquisition. Also in May 2017, we entered into a five-year senior unsecured revolving credit facility that will provide borrowing of up to \$2.25 billion when the facility becomes effective upon the closing of the Bard acquisition. This facility will expire in May 2022. Upon the effective date of the facility, it will replace the \$1.5 billion syndicated credit facility discussed further above. We will be able to issue up to \$100 million in letters of credit under this new revolving credit facility and it also includes a provision that enables BD, subject to additional commitments made by the lenders, to access up to an additional \$500 million in financing through the facility for a maximum aggregate commitment of \$2.75 billion. We will use proceeds from this facility to fund general corporate needs and to redeem, repurchase or defease certain of Bard's outstanding senior unsecured notes that will be assumed upon the closing of the acquisition.

The agreements for both the new term loan and revolving credit facility contain the following financial covenants:

- We are required to maintain an interest expense coverage ratio of not less than 4-to-1 as of the last day of each fiscal quarter. We were in compliance with this covenant relative to the term loan facility as of September 30, 2017. This covenant becomes effective for the revolving credit facility upon the effective date of the facility.
- We are required to have a leverage coverage ratio of no more than:
 - 6-to-1 from the closing date of the Bard acquisition until and including the first fiscal quarter-end thereafter;
 - 5.75-to-1 for the subsequent four fiscal quarters thereafter;
 - 5.25-to-1 for the subsequent four fiscal quarters thereafter;
 - 4.5-to-1 for the subsequent four fiscal quarters thereafter;
 - 4-to-1 for the subsequent four fiscal quarters thereafter;
 - 3.75-to-1 thereafter.

Bridge facility

Upon securing permanent financing, including the issuances of senior unsecured U.S. notes and equity securities in the third quarter of fiscal year 2017, as previously discussed above, an agreement for fully committed bridge financing of \$15.7 billion we had secured in concurrence with the announcement of the acquisition agreement was terminated.

Access to Capital and Credit Ratings

Our ability to generate cash flow from operations, issue debt, enter into other financing arrangements and attract long-term capital on acceptable terms could be adversely affected in the event there was a material decline in the demand for our products, deterioration in our key financial ratios or credit ratings, or other significantly unfavorable changes in conditions.

Our corporate credit ratings with the rating agencies Standard & Poor's Ratings Services ("S&P"), Moody's Investor Service (Moody's) and Fitch Ratings ("Fitch") were as follows at September 30, 2017:

	<u>S&P</u>	<u>Moody's</u>	<u>Fitch</u>
Ratings:			
Senior Unsecured Debt	BBB+	Baa2	BBB-
Commercial Paper	A-2	P-2	
Outlook	Negative	Negative	Stable

Upon our announcement of the agreement to acquire Bard, S&P placed our corporate credit rating of BBB+ on CreditWatch. The BBB+ rating S&P assigned to our new term loan facility and the senior unsecured U.S. notes we issued in the third quarter of fiscal year 2017 have also been placed on CreditWatch by the ratings agency. S&P has indicated that this placement will be resolved upon the acquisition's closing, which is expected to occur in the fourth calendar quarter of 2017, and that our corporate debt rating by S&P will be lowered one notch to BBB. S&P also assigned a BBB- rating to the previously discussed mandatory convertible preferred stock we issued in May 2017.

Also upon our announcement of the agreement to acquire Bard, Moody's placed our Baa2 and P-2 ratings on review for downgrade and these ratings currently remain under review. Additionally, Moody's assigned a corporate credit rating of Ba1 to our new term loan facility and to all of the tranches of senior unsecured U.S. notes issued in the third quarter, except for the tranche of 2.133% notes due June 6, 2019, which was assigned a corporate credit rating of Baa2. Moody's assigned a corporate credit rating of Baa2 to euro-denominated notes we also issued in the third quarter of fiscal year 2017. Moody's has placed the ratings assigned to the tranche of 2.133% notes and the euro-denominated notes on review for downgrade. Moody's also assigned a Ba1 rating to the BD notes we are offering in exchange of exiting Bard notes. Upon the closing of the Bard acquisition, we expect Moody's to downgrade our credit rating to below investment grade.

Additionally upon our announcement of the Bard agreement, Fitch assigned corporate debt ratings to BD for the first time and assigned BD a Long-term Issuer Default Rating of BBB- and an outlook of Stable. Fitch also assigned a BBB- rating to the euro-denominated notes we issued in the third quarter.

There were no changes to BD's long-term debt and commercial paper ratings during 2016. Lower corporate debt ratings and further downgrades of our corporate credit ratings or other credit ratings may increase our cost of borrowing. We believe that given our debt ratings, our financial management policies, our ability to generate cash flow and the non-cyclical, geographically diversified nature of our businesses, we would have access to additional short-term and long-term capital should the need arise. A rating reflects only the view of a rating agency and is not a recommendation to buy, sell or hold securities. Ratings can be revised upward or downward at any time by a rating agency if such rating agency decides that circumstances warrant such a change.

Contractual Obligations

In the normal course of business, we enter into contracts and commitments that obligate us to make payments in the future. The table below sets forth BD's significant contractual obligations and related scheduled payments as of September 30, 2017:

	Total	2018	2019 to 2020	2021 to 2022	2023 and Thereafter
	(Millions of dollars)				
Short-term debt	\$ 208	\$ 208	\$ —	\$ —	\$ —
Long-term debt(A)	26,897	615	4,986	5,001	16,295
Operating leases	277	67	104	66	39
Purchase obligations(B)	1,077	682	330	65	—
Unrecognized tax benefits(C)	—	—	—	—	—
Total(D)	<u>\$ 28,458</u>	<u>\$ 1,572</u>	<u>\$ 5,420</u>	<u>\$ 5,132</u>	<u>\$ 16,334</u>

- (A) Long-term debt obligations include expected principal and interest obligations.
- (B) Purchase obligations are for purchases made in the normal course of business to meet operational and capital requirements.
- (C) Unrecognized tax benefits at September 30, 2017 of \$349 million were all long-term in nature. Due to the uncertainty related to the timing of the reversal of these tax positions, the related liability has been excluded from the table.
- (D) Required funding obligations for 2018 relating to pension and other postretirement benefit plans are not expected to be material.

Critical Accounting Policies

The following discussion supplements the descriptions of our accounting policies contained in Note 1 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. The preparation of the consolidated financial statements requires management to use estimates and assumptions that affect the reported amounts of assets, liabilities, revenues and expenses, as well as the disclosure of contingent assets and liabilities at the date of the consolidated financial statements. Some of those judgments can be subjective and complex and, consequently, actual results could differ from those estimates. Management bases its estimates and judgments on historical experience and on various other factors that are believed to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. For any given estimate or assumption made by management, it is possible that other people applying reasonable judgment to the same facts and circumstances could develop different estimates. Actual results that differ from management's estimates could have an unfavorable effect on our consolidated financial statements. Management believes the following critical accounting policies reflect the more significant judgments and estimates used in the preparation of the consolidated financial statements:

Revenue Recognition

Some of our sales transactions qualify as multiple-element arrangements which require us to identify separate units of accounting within the arrangement and allocate the transaction consideration across these separate accounting units. For arrangements that include software and non-software elements, the transaction consideration is allocated to the software elements as a group as well as to the individual non-software elements that have been separately identified. The identification of accounting units and the allocation of total transaction consideration for multiple-element arrangements may be subjective and requires a degree of management judgment. Management's judgments relative to multiple-element arrangements may affect the timing of revenue recognition.

Transaction consideration for separately identified non-software units of accounting within an arrangement is recognized upon the completion of each deliverable based on its relative selling price. When applying the relative selling price method, the selling price of each deliverable is determined based upon the following hierarchy of evidence: vendor-specific objective evidence, which is generally based upon historical prices in stand-alone transactions; third-party evidence, which is generally based on market data on sales of similar products and services, if available; and management's best estimate of selling price. Management's best estimate of selling price is generally based upon the following considerations: stand-alone sales prices,

established price lists, costs to produce, profit margins for similar products, market conditions, and customer stratification.

For software and software-related products, we use the relative fair value method to allocate transaction consideration to each unit of accounting; whereby the evidence used in the determination of fair value estimates are based solely on vendor-specific objective evidence. To the extent that vendor specific objective evidence does not exist for delivered elements of the transaction, we apply the residual method.

The revenue allocated to equipment or instruments in multiple-element arrangements is recognized upon transfer of title and risk of loss to the customer. The revenue allocated to extended warranty contracts and software maintenance contracts is deferred and recognized as these deliverables are performed under the arrangement. The majority of deferred revenue relating to extended warranty contracts is generally recognized within a few years whereas deferred revenue relating to software maintenance contracts is generally recognized over a longer period.

Accounting for Sales-Type Leases

Our accounting for sales-type leases, which are primarily associated with dispensing products within our Medication Management Solutions unit, is based upon certain assumptions including the economic life of our leased products and the fair value of our leased products, which is used to determine the interest rate implicit to the lease. These assumptions affect that amount of gross investment in the lease, unearned income, and the sales price that is recognized relative to each sales-type lease transaction. The economic life of our leased products is based on the estimated period during which the leased product is expected to be economically usable, without limitation by the lease term, and includes an estimate of future technological advances.

The fair value of our leased products is estimated on a quarterly basis, based upon transacted cash sales prices during the preceding 12-month period, and represents normal selling price, reflecting any volume or trade discounts that may apply. Because our products are sold as part of customized systems to a diverse range of customers, many of which are affiliated with a group purchasing organization or integrated delivery network, there is a wide range of negotiated cash selling prices for our products. Accordingly, we stratify our cash selling transactions based on product configuration and customer class to determine an estimate of fair value for each product specific, within determined customer classes. Based upon this stratification, we calculate the weighted average selling price of each configured product using the interquartile range methodology and remove outliers from the population of normal cash selling prices, which narrows the range of selling prices within each stratified customer class. The resulting weighted average selling price is the single point estimate of fair value that we use as the normal selling price and this fair value estimate is used to determine the implicit interest rate for each product subject to a sales-type lease arrangement. In certain instances, we do not have sufficient corresponding historical cash selling transactions to support fair values of specific combinations of product configurations and customer classes. In these instances, fair value is estimated by applying the average discount percentage given to the respective customer class, over the trailing 12 months, to the list price of the products whose fair value was not determined using the interquartile range methodology described above. The resulting fair value(s) is then used to derive the implicit rate of the lease. The interest rate implicit to the lease is then used to determine the amount of revenue recognized at the inception of the lease and the revenue recognized over the life of the lease.

Our net investment in sales-type leases primarily arises from the leasing of dispensing equipment and as such, the methodology for determining the relating allowance for credit losses is based on the collective population and is not stratified by class or portfolio segment. The allowance for credit losses is based on historical experience loss rates as well as on management's judgments regarding the potential impact of anticipated changes in business practices, market dynamics, and economic conditions. These assumptions are inherently subjective and it is possible that we will experience actual credit losses that are different from our current estimates.

In April 2017, in conjunction with the implementation of a new go-to-market business model for our dispensing business, we amended new and existing leases to provide a limited return option. Prior to the amendment these leases were accounted for as sales-type leases in accordance with Accounting Standards

Codification 840, Leases, as the typical non-cancellable lease term of 5 years exceeded 75% of the equipment's estimated useful life and the present value of the minimum lease payments exceeded 90% of the equipment's fair value. As sales-type leases, we recognized revenue upon installation of equipment at a customer site based on the present value of the minimum lease payments. As a result of the contract amendment, the amended lease term is shortened and as a result, the majority of leases no longer meet the criteria for recognition as sales-type leases. Accordingly, the leases have been classified as operating leases as of the modification date and revenue is generally recognized ratably over the lease term.

Accounting for Software Products

We sell and lease products with embedded software and as such, we must evaluate these products to determine if industry-specific revenue recognition requirements apply to these sales transactions. This evaluation process is often complex and subject to significant judgment. If software is considered not essential to the non-software elements of a product but is considered more than incidental to a product as a whole, the product's software elements must be separated from its non-software elements under the requirements relating to multiple-element arrangements. The product's software elements must be accounted for under software industry-specific revenue recognition requirements and the application of these requirements may significantly affect the timing and amount of revenue recognized.

While we have determined that the software embedded in the following product groupings is more than incidental to the products as a whole, the non-software elements (i.e., hardware) and software elements work together to deliver the essential functionality of these products as a whole. As such, the accounting for these product offerings does not fall within the scope of software industry-specific accounting requirements:

- Infusion products (when sold with safety software, patient identification products and certain diagnostic equipment) within our Medication Management Solutions unit;
- Dispensing products within our Medication Management Solutions unit;
- Research and clinical instruments within our Biosciences unit.

Our standalone software application sales and any related post-contract support related to these sales are accounted for under the software industry-specific revenue recognition requirements.

Impairment of Assets

Goodwill and in-process research and development assets are subject to impairment reviews at least annually, or whenever indicators of impairment arise. Intangible assets with finite lives, including developed technology, and other long-lived assets, are periodically reviewed for impairment when impairment indicators are present.

We assess goodwill for impairment at the reporting unit level, which is defined as an operating segment or one level below an operating segment, referred to as a component. Our reporting units generally represent one level below reporting segments and we aggregate components within an operating segment that have similar economic characteristics. Potential impairment of goodwill is identified by comparing the fair value of a reporting unit with its carrying value. Our annual goodwill impairment test performed on July 1, 2017 did not result in any impairment charges, as the fair value of each reporting unit exceeded its carrying value.

We generally use the income approach to derive the fair value for impairment assessments. This approach calculates fair value by estimating future cash flows attributable to the assets and then discounting these cash flows to a present value using a risk-adjusted discount rate. We selected this method because we believe the income approach most appropriately measures our income producing assets. This approach requires significant management judgment with respect to future volume, revenue and expense growth rates, changes in working capital use, appropriate discount rates and other assumptions and estimates. The estimates and assumptions used are consistent with BD's business plans. The use of alternative estimates and assumptions could increase or decrease the estimated fair value of the asset, and potentially result in different impacts to BD's results of operations. Actual results may differ from management's estimates.

Income Taxes

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

BD conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of tax jurisdictions. In evaluating the exposure associated with various tax filing positions, we record accruals for uncertain tax positions based on the technical support for the positions, our past audit experience with similar situations, and the potential interest and penalties related to the matters. BD's effective tax rate in any given period could be impacted if, upon resolution with taxing authorities, we prevailed in positions for which reserves have been established, or we were required to pay amounts in excess of established reserves.

BD has reviewed its needs in the U.S. for possible repatriation of undistributed earnings of its foreign subsidiaries and, with exception for certain countries, continues to invest foreign subsidiaries earnings outside of the U.S. to fund foreign investments or meet foreign working capital and property, plant and equipment expenditure needs. Deferred taxes are not provided on undistributed earnings of foreign subsidiaries that are indefinitely reinvested. At September 30, 2017, the cumulative amount of such undistributed earnings indefinitely reinvested outside the United States was \$9.6 billion. The determination of the amount of the unrecognized deferred tax liability related to the undistributed earnings is not practicable because of the complexities associated with its hypothetical calculation.

Contingencies

We are involved, both as a plaintiff and a defendant, in various legal proceedings that arise in the ordinary course of business, including, without limitation, product liability, antitrust and environmental matters, as further discussed in Note 5 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. We assess the likelihood of any adverse judgments or outcomes to these matters as well as potential ranges of probable losses. We establish accruals to the extent probable future losses are estimable (in the case of environmental matters, without considering possible third-party recoveries). A determination of the amount of accruals, if any, for these contingencies is made after careful analysis of each individual issue and, when appropriate, is developed after consultation with outside counsel. The accruals may change in the future due to new developments in each matter or changes in our strategy in dealing with these matters.

Given the uncertain nature of litigation generally, we are not able in all cases to estimate the amount or range of loss that could result from an unfavorable outcome of the litigation to which we are a party. In view of these uncertainties, we could incur charges in excess of any currently established accruals and, to the extent available, excess liability insurance. In the opinion of management, any such future charges, individually or in the aggregate, could have a material adverse effect on BD's consolidated results of operations and consolidated net cash flows.

Benefit Plans

We have significant net pension and other postretirement and postemployment benefit costs that are measured using actuarial valuations. These benefit costs include assumptions for the discount rate. Pension benefit costs also include an assumption for the expected return on plan assets. These assumptions have a significant effect on the amounts reported. In addition to the analysis below, see Note 8 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data for additional discussion.

The discount rate is selected each year based on investment grade bonds and other factors as of the measurement date (September 30). Specifically for the U.S. pension plan, we will use a discount rate of 3.73% for 2018, which was based on an actuarially-determined, company-specific yield curve to measure liabilities as

of the measurement date. To calculate the pension expense in 2018, we will apply the individual spot rates along the yield curve that correspond with the timing of each future cash outflow for benefit payments in order to calculate interest cost and service cost. Additional disclosures regarding the method to be used in calculating the interest cost and service cost components of pension expense for 2018 are provided in Note 8 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data. The expected long-term rate of return on plan assets assumption, although reviewed each year, changes less frequently due to the long-term nature of the assumption. This assumption does not impact the measurement of assets or liabilities as of the measurement date; rather, it is used only in the calculation of pension expense. To determine the expected long-term rate of return on pension plan assets, we consider many factors, including our historical assumptions compared with actual results; benchmark data; expected returns on various plan asset classes, as well as current and expected asset allocations. We will use a long-term expected rate of return on plan assets assumption of 7.25% for the U.S. pension plan in 2018. We believe our discount rate and expected long-term rate of return on plan assets assumptions are appropriate based upon the above factors.

Sensitivity to changes in key assumptions for our U.S. pension and other postretirement and postemployment plans are as follows:

- Discount rate — A change of plus (minus) 25 basis points, with other assumptions held constant, would have an estimated \$5 million favorable (unfavorable) impact on the total U.S. net pension and other postretirement and postemployment benefit plan costs. This estimate assumes no change in the shape or steepness of the company-specific yield curve used to plot the individual spot rates that will be applied to the future cash outflows for future benefit payments in order to calculate interest and service cost.
- Expected return on plan assets — A change of plus (minus) 25 basis points, with other assumptions held constant, would have an estimated \$3 million favorable (unfavorable) impact on U.S. pension plan costs.

Share-Based Compensation

Compensation cost relating to share-based payment transactions is recognized in net income using a fair value measurement method. All share-based payments to employees, including grants of employee stock options, are recognized in the statement of operations as compensation expense (based on their fair values) over the vesting period of the awards. We determine the fair value of certain share-based awards using a lattice-based binomial option valuation model that incorporates certain assumptions, such as the risk-free interest rate, expected volatility, expected dividend yield and expected life of the options. See Note 7 to the consolidated financial statements contained in Item 8. Financial Statements and Supplementary Data for additional discussion.

Cautionary Statement Regarding Forward-Looking Statements

BD and its representatives may from time to time make certain forward-looking statements in publicly released materials, both written and oral, including statements contained in filings with the Securities and Exchange Commission, press releases, and our reports to shareholders. Forward-looking statements may be identified by the use of words such as “plan,” “expect,” “believe,” “intend,” “will,” “may”, “anticipate,” “estimate” and other words of similar meaning in conjunction with, among other things, discussions of future operations and financial performance (including volume growth, sales and earnings per share growth, and cash flows) and statements regarding our strategy for growth, future product development, regulatory approvals, competitive position and expenditures. All statements that address our future operating performance or events or developments that we expect or anticipate will occur in the future are forward-looking statements.

Forward-looking statements are, and will be, based on management’s then-current views and assumptions regarding future events, developments and operating performance, and speak only as of their dates. Investors should realize that if underlying assumptions prove inaccurate, or risks or uncertainties materialize, actual results could vary materially from our expectations and projections. Investors are therefore cautioned not to place undue reliance on any forward-looking statements. Furthermore, we undertake no obligation to update or revise any forward-looking statements after the date they are made, whether as a result of new information,

future events and developments or otherwise, except as required by applicable law or regulations.

The following are some important factors that could cause our actual results to differ from our expectations in any forward-looking statements. For further discussion of certain of these factors, see Item 1A. Risk Factors.

- Weakness in the global economy and financial markets, which could increase the cost of operating our business, weaken demand for our products and services, negatively impact the prices we can charge for our products and services, or impair our ability to produce our products.
- Competitive factors that could adversely affect our operations, including new product introductions (for example, new forms of drug delivery) by our current or future competitors, increased pricing pressure due to the impact of low-cost manufacturers, patents attained by competitors (particularly as patents on our products expire), and new entrants into our markets.
- The adverse financial impact resulting from unfavorable changes in foreign currency exchange rates.
- Regional, national and foreign economic factors, including inflation, deflation, and fluctuations in interest rates, and their potential effect on our operating performance.
- Our ability to achieve our projected level or mix of product sales, as our earnings forecasts are based on projected sales volumes and pricing of many product types, some of which are more profitable than others.
- Changes in reimbursement practices of third-party payers or adverse decisions relating to our products by such payers, which could reduce demand for our products or the price we can charge for such products.
- The impact of the medical device excise tax under the Patient Protection and Affordable Care Act (the "PPACA") in the United States. This tax has been suspended through December 31, 2017, and it is uncertain whether the suspension will be extended beyond that date.
- Healthcare reform in the U.S. or in other countries in which we do business that may involve changes in government pricing and reimbursement policies or other cost containment reforms.
- Changes in domestic and foreign healthcare industry practices that result in a reduction in procedures using our products or increased pricing pressures, including the continued consolidation among healthcare providers and trends toward managed care and healthcare cost containment.
- The impact of changes in U.S. federal laws and policy that could affect fiscal and tax policies, healthcare, and international trade agreements.
- Fluctuations in the cost and availability of oil-based resins and other raw materials, as well as certain components, used in our products, the ability to maintain favorable supplier arrangements and relationships (particularly with respect to sole-source suppliers), and the potential adverse effects of any disruption in the availability of such items.
- Security breaches of our information technology systems or our products, which could impair our ability to conduct business, result in the loss of BD trade secrets or otherwise compromise sensitive information of BD or its customers, suppliers and other business partners, or of customers' patients, or result in product efficacy or safety concerns for certain of our products.
- Difficulties inherent in product development, including the potential inability to successfully continue technological innovation, successfully complete clinical trials, obtain regulatory approvals in the United States and abroad, obtain intellectual property protection for our products, obtain coverage and adequate reimbursement for new products, or gain and maintain market approval of products, as well as the possibility of infringement claims by competitors with respect to patents or other intellectual property rights, all of which can preclude or delay commercialization of a product. Delays in obtaining necessary approvals or clearances from the FDA or other regulatory agencies or changes in the

regulatory process may also delay product launches and increase development costs.

- The impact of business combinations, including any volatility in earnings relating to acquisition-related costs, and our ability to successfully integrate any business we may acquire.
- Our ability to penetrate or expand our operations in emerging markets, which depends on local economic and political conditions, and how well we are able to acquire or form strategic business alliances with local companies and make necessary infrastructure enhancements to production facilities and distribution networks. Our international operations also increase our compliance risks, including risks under the Foreign Corrupt Practices Act and other anti-corruption laws.
- Political conditions in international markets, including civil unrest, terrorist activity, governmental changes, trade barriers, restrictions on the ability to transfer capital across borders and governmental expropriation of assets. This includes the possible impact of the June 2016 advisory referendum by British voters to exit the European Union, which has created uncertainties affecting business operations in the United Kingdom and the EU.
- Deficit reduction efforts or other actions that reduce the availability of government funding for healthcare and research, which could weaken demand for our products and result in additional pricing pressures, as well as create potential collection risks associated with such sales.
- Fluctuations in university or U.S. and international governmental funding and policies for life sciences research.
- Fluctuations in the demand for products we sell to pharmaceutical companies that are used to manufacture, or are sold with, the products of such companies, as a result of funding constraints, consolidation or otherwise.
- The effects of events that adversely impact our ability to manufacture our products (particularly where production of a product line is concentrated in one or more plants) or our ability to source materials or components from suppliers (including sole-source suppliers) that are needed for such manufacturing. In particular, damage to our manufacturing facilities in Puerto Rico resulting from Hurricane Maria in September 2017 could adversely impact our revenue and earnings results for fiscal year 2018.
- Pending and potential future litigation or other proceedings adverse to BD, including antitrust, product liability, environmental and patent infringement, and the availability or collectability of insurance relating to any such claims.
- New or changing laws and regulations affecting our domestic and foreign operations, or changes in enforcement practices, including laws relating to trade, monetary and fiscal policies, taxation (including tax reforms that could adversely impact multinational corporations), sales practices, environmental protection, price controls, and licensing and regulatory requirements for new products and products in the postmarketing phase. In particular, the U.S. and other countries may impose new requirements regarding registration, labeling or prohibited materials that may require us to re-register products already on the market or otherwise impact our ability to market our products. Environmental laws, particularly with respect to the emission of greenhouse gases, are also becoming more stringent throughout the world, which may increase our costs of operations or necessitate changes in our manufacturing plants or processes or those of our suppliers, or result in liability to BD.
- Product efficacy or safety concerns regarding our products resulting in product recalls, regulatory action on the part of the U.S. Food and Drug Administration (FDA) or foreign counterparts, declining sales and product liability claims, and damage to our reputation. As a result of the CareFusion acquisition, we are operating under a consent decree with the FDA relating to our U.S. infusion pump business. The consent decree authorizes the FDA, in the event of any violations in the future, to order us to cease manufacturing and distributing products, recall products or take other actions, and we may be required to pay significant monetary damages if we fail to comply with any provision of the consent decree.
- Risks relating to our acquisition of CareFusion, including our ability to continue to successfully

combine and integrate the CareFusion operations in order to fully obtain the anticipated benefits and costs savings from the transaction.

- Risks related to our pending acquisition of Bard, including:
 - The failure to satisfy the conditions to completing the transaction, including obtaining required regulatory approvals.
 - Conditions to obtaining regulatory approval that may place restrictions on the business of the combined company.
 - Our failure to obtain the anticipated benefits and costs savings from the acquisition.
 - The impact of the additional debt we incurred and the equity and equity-linked securities that we issued to finance the acquisition, including on our credit ratings and costs of borrowing.
- The effect of adverse media exposure or other publicity regarding BD's business or operations, including the effect on BD's reputation or demand for its products.
- The effect of market fluctuations on the value of assets in BD's pension plans and on actuarial interest rate and asset return assumptions, which could require BD to make additional contributions to the plans or increase our pension plan expense.
- Our ability to obtain the anticipated benefits of restructuring programs, if any, that we may undertake.
- Issuance of new or revised accounting standards by the Financial Accounting Standards Board or the Securities and Exchange Commission.

The foregoing list sets forth many, but not all, of the factors that could impact our ability to achieve results described in any forward-looking statements. Investors should understand that it is not possible to predict or identify all such factors and should not consider this list to be a complete statement of all potential risks and uncertainties.

Item 7A. *Quantitative and Qualitative Disclosures About Market Risk.*

The information required by this item is included in Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations, and in Notes 1, 13 and 14 to the consolidated financial statements contained in Item 8, Financial Statements and Supplementary Data, and is incorporated herein by reference.

Item 8. Financial Statements and Supplementary Data.

Reports of Management

Management's Responsibilities

The following financial statements have been prepared by management in conformity with U.S. generally accepted accounting principles and include, where required, amounts based on the best estimates and judgments of management. The integrity and objectivity of data in the financial statements and elsewhere in this Annual Report are the responsibility of management.

In fulfilling its responsibilities for the integrity of the data presented and to safeguard the Company's assets, management employs a system of internal accounting controls designed to provide reasonable assurance, at appropriate cost, that the Company's assets are protected and that transactions are appropriately authorized, recorded and summarized. This system of control is supported by the selection of qualified personnel, by organizational assignments that provide appropriate delegation of authority and division of responsibilities, and by the dissemination of written policies and procedures. This control structure is further reinforced by a program of internal audits, including a policy that requires responsive action by management.

The Board of Directors monitors the internal control system, including internal accounting and financial reporting controls, through its Audit Committee, which consists of eight independent Directors. The Audit Committee meets periodically with the independent registered public accounting firm, the internal auditors and management to review the work of each and to satisfy itself that they are properly discharging their responsibilities. The independent registered public accounting firm and the internal auditors have full and free access to the Audit Committee and meet with its members, with and without management present, to discuss the scope and results of their audits including internal control, auditing and financial reporting matters.

Management's Report on Internal Control Over Financial Reporting

Management is responsible for establishing and maintaining adequate internal control over financial reporting, as defined in Rule 13a-15(f) under the Securities Act of 1934. Management conducted an assessment of the effectiveness of internal control over financial reporting based on the criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework). Based on this assessment and those criteria, management concluded that internal control over financial reporting was effective as of September 30, 2017.

The financial statements and internal control over financial reporting have been audited by Ernst & Young LLP, an independent registered public accounting firm. Ernst & Young's reports with respect to fairness of the presentation of the financial statements, and the effectiveness of internal control over financial reporting, are included herein.

/s/ Vincent A. Forlenza

Vincent A. Forlenza

Chairman and Chief Executive Officer

/s/ Christopher Reidy

Christopher Reidy

*Executive Vice President, Chief
Financial Officer and Chief
Administrative Officer*

/s/ John Gallagher

John Gallagher

*Senior Vice President, Corporate
Finance, Controller and Treasurer*

Report of Independent Registered Public Accounting Firm

To the Shareholders and Board of Directors of
Becton, Dickinson and Company

We have audited the accompanying consolidated balance sheets of Becton, Dickinson and Company as of September 30, 2017 and 2016, and the related consolidated statements of income, comprehensive income and cash flows for each of the three years in the period ended September 30, 2017. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

We conducted our audits in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the consolidated financial position of Becton, Dickinson and Company at September 30, 2017 and 2016, and the consolidated results of its operations and its cash flows for each of the three years in the period ended September 30, 2017, in conformity with U.S. generally accepted accounting principles.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States), Becton, Dickinson and Company's internal control over financial reporting as of September 30, 2017, based on criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework) and our report dated November 22, 2017 expressed an unqualified opinion thereon.

/s/ ERNST & YOUNG LLP

New York, New York
November 22, 2017

Report of Independent Registered Public Accounting Firm

To the Shareholders and Board of Directors of
Becton, Dickinson and Company

We have audited Becton, Dickinson and Company's internal control over financial reporting as of September 30, 2017, based on criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework) (the COSO criteria). Becton, Dickinson and Company's management is responsible for maintaining effective internal control over financial reporting, and for its assessment of the effectiveness of internal control over financial reporting included in the accompanying Management's Report on Internal Control Over Financial Reporting. Our responsibility is to express an opinion on the company's internal control over financial reporting based on our audit.

We conducted our audit in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether effective internal control over financial reporting was maintained in all material respects. Our audit included obtaining an understanding of internal control over financial reporting, assessing the risk that a material weakness exists, testing and evaluating the design and operating effectiveness of internal control based on the assessed risk, and performing such other procedures as we considered necessary in the circumstances. We believe that our audit provides a reasonable basis for our opinion.

A company's internal control over financial reporting is a process designed 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. A company's internal control over financial reporting includes those policies and procedures that (1) pertain to the maintenance of records that, in reasonable detail, accurately and fairly reflect the transactions and dispositions of the assets of the company; (2) provide reasonable assurance that transactions are recorded as necessary to permit preparation of financial statements in accordance with generally accepted accounting principles, and that receipts and expenditures of the company are being made only in accordance with authorizations of management and directors of the company; and (3) provide reasonable assurance regarding prevention or timely detection of unauthorized acquisition, use, or disposition of the company's assets that could have a material effect on the financial statements.

Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. Also, projections of any evaluation of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

In our opinion, Becton, Dickinson and Company maintained, in all material respects, effective internal control over financial reporting as of September 30, 2017, based on the COSO criteria.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States), the consolidated balance sheets of Becton, Dickinson and Company as of September 30, 2017 and 2016, and the related consolidated statements of income, comprehensive income and cash flows for each of the three years in the period ended September 30, 2017 of Becton, Dickinson and Company, and our report dated November 22, 2017 expressed an unqualified opinion thereon.

/s/ ERNST & YOUNG LLP

New York, New York
November 22, 2017

Consolidated Statements of Income

Becton, Dickinson and Company Years Ended September 30

Millions of dollars, except per share amounts

	2017	2016	2015
Revenues	\$ 12,093	\$ 12,483	\$ 10,282
Cost of products sold	6,151	6,492	5,587
Selling and administrative expense	2,925	3,005	2,563
Research and development expense	774	828	632
Acquisitions and other restructurings	354	728	426
Other operating expense, net	410	—	—
Total Operating Costs and Expenses	<u>10,615</u>	<u>11,053</u>	<u>9,207</u>
Operating Income	1,478	1,430	1,074
Interest expense	(521)	(388)	(371)
Interest income	76	21	15
Other (expense) income, net	(57)	11	21
Income Before Income Taxes	<u>976</u>	<u>1,074</u>	<u>739</u>
Income tax (benefit) provision	(124)	97	44
Net Income	<u>1,100</u>	<u>976</u>	<u>695</u>
Preferred stock dividends	(70)	—	—
Net income applicable to common shareholders	<u>\$ 1,030</u>	<u>\$ 976</u>	<u>\$ 695</u>
Basic Earnings per Share	<u>\$ 4.70</u>	<u>\$ 4.59</u>	<u>\$ 3.43</u>
Diluted Earnings per Share	<u>\$ 4.60</u>	<u>\$ 4.49</u>	<u>\$ 3.35</u>

Consolidated Statements of Comprehensive Income

Becton, Dickinson and Company Years Ended September 30

Millions of dollars	2017	2016	2015
Net Income	\$ 1,100	\$ 976	\$ 695
Other Comprehensive Income (Loss), Net of Tax			
Foreign currency translation adjustments	11	(50)	(692)
Defined benefit pension and postretirement plans	179	(141)	(36)
Cash flow hedges	17	1	(9)
Other Comprehensive Income (Loss), Net of Tax	206	(191)	(737)
Comprehensive Income (Loss)	\$ 1,306	\$ 786	\$ (42)

Consolidated Balance Sheets
Becton, Dickinson and Company
September 30

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

	<u>2017</u>	<u>2016</u>
Assets		
Current Assets		
Cash and equivalents	\$ 14,179	\$ 1,541
Short-term investments	21	27
Trade receivables, net	1,744	1,618
Current portion of net investment in sales-type leases	16	339
Inventories	1,818	1,719
Assets held for sale	—	642
Prepaid expenses and other	856	480
Total Current Assets	<u>18,633</u>	<u>6,367</u>
Property, Plant and Equipment, Net	4,638	3,901
Goodwill	7,563	7,419
Customer Relationships, Net	2,830	3,022
Developed Technology, Net	2,478	2,655
Other Intangibles, Net	585	604
Net Investment in Sales-Type Leases, Less Current Portion	38	796
Other Assets	968	824
Total Assets	<u>\$ 37,734</u>	<u>\$ 25,586</u>
Liabilities and Shareholders' Equity		
Current Liabilities		
Short-term debt	\$ 203	\$ 1,001
Accounts payable	797	665
Accrued expenses	1,393	1,575
Salaries, wages and related items	773	696
Income taxes	176	274
Liabilities held for sale	—	189
Total Current Liabilities	<u>3,342</u>	<u>4,400</u>
Long-Term Debt	18,667	10,550
Long-Term Employee Benefit Obligations	1,168	1,319
Deferred Income Taxes and Other	1,609	1,684
Commitments and Contingencies (See Note 5)		
Shareholders' Equity		
Preferred stock	2	—
Common stock — \$1 par value: authorized — 640,000,000 shares; issued — 346,687,160 shares in 2017 and 332,662,160 shares in 2016.	347	333
Capital in excess of par value	9,619	4,693
Retained earnings	13,111	12,727
Deferred compensation	19	22
Common stock in treasury — at cost — 118,744,758 shares in 2017 and 119,370,934 shares in 2016.	(8,427)	(8,212)
Accumulated other comprehensive loss	(1,723)	(1,929)
Total Shareholders' Equity	<u>12,948</u>	<u>7,633</u>
Total Liabilities and Shareholders' Equity	<u>\$ 37,734</u>	<u>\$ 25,586</u>

Consolidated Statements of Cash Flows

Becton, Dickinson and Company Years Ended September 30

Millions of dollars	2017	2016	2015
Operating Activities			
Net income	\$ 1,100	\$ 976	\$ 695
Adjustments to net income to derive net cash provided by operating activities:			
Depreciation and amortization	1,088	1,114	891
Share-based compensation	174	196	166
Deferred income taxes	(236)	(426)	(336)
Change in operating assets and liabilities:			
Trade receivables, net	(93)	(128)	(2)
Net investment in sales-type leases	14	51	28
Inventories	(46)	69	200
Prepaid expenses and other	(380)	39	(97)
Accounts payable, income taxes and other liabilities	134	368	145
Pension obligation	84	(32)	28
Excess tax benefits from payments under share-based compensation plans	77	—	—
Lease contract modification-related charge	748	—	—
Other, net	(114)	332	11
Net Cash Provided by Operating Activities	<u>2,550</u>	<u>2,559</u>	<u>1,730</u>
Investing Activities			
Capital expenditures	(727)	(693)	(596)
Proceeds from (purchases of) investments, net	13	(1)	840
Acquisitions of businesses, net of cash acquired	(174)	—	(8,414)
Divestitures of businesses, net	165	158	—
Other, net	(161)	(133)	(147)
Net Cash Used for Investing Activities	<u>(883)</u>	<u>(669)</u>	<u>(8,318)</u>
Financing Activities			
Change in short-term debt	(200)	(500)	497
Proceeds from long-term debt	11,462	—	6,164
Payments of debt	(3,980)	(752)	(6)
Proceeds from issuance of equity securities	4,827	—	—
Repurchase of common stock	(220)	—	—
Excess tax benefit from payments under share-based compensation plans	—	86	48
Dividends paid	(677)	(562)	(485)
Other, net	(234)	(32)	(27)
Net Cash Provided by (Used for) Financing Activities	<u>10,977</u>	<u>(1,761)</u>	<u>6,190</u>
Effect of exchange rate changes on cash and equivalents	(6)	(12)	(38)
Net Increase (Decrease) in Cash and Equivalents	<u>12,638</u>	<u>117</u>	<u>(436)</u>
Opening Cash and Equivalents	1,541	1,424	1,861
Closing Cash and Equivalents	<u>\$ 14,179</u>	<u>\$ 1,541</u>	<u>\$ 1,424</u>

Notes to Consolidated Financial Statements
Becton, Dickinson and Company
Millions of dollars, except per share amounts and numbers of shares

Note 1 — Summary of Significant Accounting Policies

Basis of Presentation

The accompanying Consolidated Financial Statements and Notes to Consolidated Financial Statements of Becton, Dickinson and Company (the “Company”) have been prepared in accordance with U.S. generally accepted accounting principles. Within the financial statements and tables presented, certain columns and rows may not add due to the use of rounded numbers for disclosure purposes. Percentages and earnings per share amounts presented are calculated from the underlying amounts. Our fiscal year ends on September 30.

Principles of Consolidation

The consolidated financial statements include the Company’s accounts and those of its majority-owned subsidiaries after the elimination of intercompany transactions. The Company has no material interests in variable interest entities.

Cash Equivalents

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

Short-Term Investments

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

Trade and Financing Receivables

The Company grants credit to customers in the normal course of business and the resulting trade receivables are stated at their net realizable value. The allowance for doubtful accounts represents the Company’s estimate of probable credit losses relating to trade receivables and is determined based on historical experience and other specific account data. Amounts are written off against the allowances for doubtful accounts when the Company determines that a customer account is uncollectible.

Inventories

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

Property, Plant and Equipment

Property, plant and equipment are stated at cost, less accumulated depreciation and amortization. Depreciation and amortization are principally provided on the straight-line basis over estimated useful lives, which range from 20 to 45 years for buildings, four to 13 years for machinery and equipment and one to 12 years for leasehold improvements. Depreciation and amortization expense was \$406 million, \$452 million and \$417 million in fiscal years 2017, 2016 and 2015, respectively.

Goodwill and Other Intangible Assets

The Company’s unamortized intangible assets include goodwill and in-process research and development assets which arise from acquisitions. The Company currently reviews all indefinite-lived assets, including goodwill, for impairment using quantitative models. Goodwill is reviewed at least annually for impairment at the reporting unit level, which is defined as an operating segment or one level below an operating segment, referred to as a component. The Company’s reporting units generally represent one level below reporting segments, and components within an operating segment that have similar economic characteristics are

aggregated. Potential impairment of goodwill is identified by comparing the fair value of a reporting unit, estimated using an income approach, with its carrying value. The annual impairment review performed on July 1, 2017 indicated that all identified reporting units' fair values exceeded their respective carrying values.

The review for impairment of in-process research and development assets is performed by comparing the fair value of the technology or project assets, estimated using an income approach, with their carrying value. In-process research and development assets are considered indefinite-lived assets and are reviewed at least annually for impairment until projects are completed or abandoned. Certain trademarks that are considered to generate cash flows indefinitely are also considered to be indefinite-lived intangible assets and these assets are also reviewed at least annually for impairment.

Amortized intangible assets include developed technology assets which arise from acquisitions. These assets represent acquired intellectual property that is already technologically feasible upon the acquisition date or acquired in-process research and development assets that are completed subsequent to acquisition. Developed technology assets are generally amortized over periods ranging from 15 to 20 years, using the straight-line method. Other intangibles with finite useful lives, which include patents, are amortized over periods principally ranging from one to 40 years, using the straight-line method. Finite-lived intangible assets, including developed technology assets, are periodically reviewed when impairment indicators are present to assess recoverability from future operations using undiscounted cash flows. The carrying values of these finite-lived assets are compared to the undiscounted cash flows they are expected to generate and an impairment loss is recognized in operating results to the extent any finite-lived intangible asset's carrying value exceeds its calculated fair value.

Foreign Currency Translation

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

Revenue Recognition

Revenue from product sales is typically recognized when all of the following criteria have been met: persuasive evidence of an arrangement exists; delivery has occurred or services have been rendered; product price is fixed or determinable; collection of the resulting receivable is reasonably assured. Certain sales arrangements contain multiple deliverables, including equipment and service deliverables, which requires the Company to determine the separate units of account. If the deliverable meets the criteria of a separate unit of accounting, the arrangement consideration is allocated to each element based upon its relative selling price. In determining the best evidence of selling price of a unit of account the Company utilizes vendor-specific objective evidence ("VSOE"), which is the price the Company charges when the deliverable is sold separately. When VSOE is not available, management uses relevant third-party evidence ("TPE") of selling price, if available. When neither VSOE nor TPE of selling price exists, management uses its best estimate of selling price.

Revenue allocated to equipment deliverables is recognized upon customer acceptance, which occurs after the transfer of title and risk of loss to the customer and the completion of installation or training services. When related training services are considered inconsequential, delivery is deemed to occur upon the transfer of title and risk of loss, at which time revenue and the costs associated with installation and training are recognized.

For equipment lease revenue, transactions are evaluated and classified as either operating leases or sales-type leases. Lease income for products sold under sales-type leases is recognized as revenue upon the completion of installation activities in the amount of the present value of the minimum lease payments. The financing component of sales-type leases is recorded as revenue over the lease term. For products sold under operating leases, revenue is recognized at the contracted rate over the rental period, as defined within the customer agreement.

For products sold and leased with embedded software, if software is considered not essential to the non-software elements of a product but is considered more than incidental to a product as a whole, the product's software elements must be separated from its non-software elements under the requirements relating to multiple-element arrangements and accounted for under software industry-specific revenue recognition requirements. However, if it is determined that the embedded software is more than incidental to the product as a whole but the non-software elements and software elements work together to deliver the essential functionality of the products as a whole, then the accounting for such product does not fall within the scope of software industry-specific accounting requirements.

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

Shipping and Handling Costs

Shipping and handling costs are included in *Selling and administrative expense*. Shipping expense was \$365 million, \$401 million and \$351 million in 2017, 2016 and 2015, respectively.

Derivative Financial Instruments

All derivatives are recorded in the balance sheet at fair value and changes in fair value are recognized currently in earnings unless specific hedge accounting criteria are met. Any deferred gains or losses associated with derivative instruments are recognized in income in the period in which the underlying hedged transaction is recognized. Additional disclosures regarding the Company's accounting for derivative instruments are provided in Note 13.

Income Taxes

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

The Company conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of tax jurisdictions. In evaluating the exposure associated with various tax filing positions, the Company records accruals for uncertain tax positions, based on the technical support for the positions, past audit experience with similar situations, and the potential interest and penalties related to the matters.

The Company maintains valuation allowances where it is more likely than not that all or a portion of a deferred tax asset will not be realized. Changes in valuation allowances are included in the tax provision in the period of change. In determining whether a valuation allowance is warranted, management evaluates factors such as prior earnings history, expected future earnings, carryback and carryforward periods and tax strategies that could potentially enhance the likelihood of the realization of a deferred tax asset. Additional disclosures regarding the Company's accounting for income taxes are provided in Note 16.

Earnings per Share

Basic earnings per share are computed by dividing income available to common stockholders by the weighted average number of common shares outstanding. Diluted earnings per share reflect the potential dilution that could occur if securities or other contracts to issue common stock were exercised or converted into common stock. In computing diluted earnings per share, only potential common shares that are dilutive (i.e., those that reduce earnings per share or increase loss per share) are included in the calculation.

Use of Estimates

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

Note 2 — Accounting Changes

New Accounting Principle Adopted

On October 1, 2016, the Company prospectively adopted amended requirements issued by the Financial Accounting Standards Board ("FASB") relating to the timing of recognition and classification of share-based compensation award-related income tax effects. Upon the settlement of awards during fiscal year 2017, the Company recorded tax benefits for the year ended of \$77 million to *Income tax provision (benefit)* within its consolidated statement of income. These tax benefits were recorded within *Capital in excess of par value* on the Company's consolidated balance sheet in the prior-year period. Because these excess tax benefits are no longer recorded in *Capital in excess of par value*, the current period adjustment for the dilutive impact of share equivalents from share-based plans, which is used in the Company's computation of diluted earnings per share, increased by approximately 1 million shares. Also per the amended guidance, the Company classified the \$77 million of excess tax benefits for the year ended September 30, 2017 on its consolidated statement of cash flows within *Net Cash Provided by Operating Activities*, rather than *Net Cash Provided by (Used for) Financing Activities*, which included the excess tax benefits for the years ended September 30, 2016 and 2015. The amended guidance allows entities to account for award forfeitures as they occur; however, the Company has elected to continue its determination of compensation cost recognized in each period based upon an estimate of expected future forfeitures.

New Accounting Principles Not Yet Adopted

In February 2016, the FASB issued a new lease accounting standard which requires lessees to recognize lease assets and lease liabilities on the balance sheet. The new standard also requires expanded disclosures regarding leasing arrangements. The Company is currently evaluating the impact that this new lease accounting standard will have on its consolidated financial statements upon its adoption of the standard on October 1, 2019.

In May 2014, the FASB issued a new revenue recognition standard. Under this standard, revenue will be recognized upon the transfer of goods or services to customers and the amount of revenue recognized will reflect the consideration to which a reporting entity expects to be entitled in exchange for those goods or services. The Company intends to adopt the standard, as required, on October 1, 2018 and recently completed an initial assessment to identify the potential areas of impact that this new revenue recognition standard will have on its consolidated financial statements. As part of the initial assessment, the Company reviewed a representative sample of its contracts across its various businesses and geographies to identify potential differences that could result from applying the requirements of the new standard. The analysis included identifying whether there may be differences in timing of revenue recognition under the new standard as well as assessing performance obligations, variable consideration, and contract costs. The Company has not yet estimated the impact of the new standard on the timing and pattern of its revenue recognition. The Company continues to evaluate the available adoption methods, and apprises its audit committee of the project status regularly.

Note 3 — Shareholders' Equity

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

(Millions of dollars)	Common Stock Issued at Par Value	Capital in Excess of Par Value	Retained Earnings	Deferred Compensation	Treasury Stock	
					Shares (in thousands)	Amount
Balance at September 30, 2014	\$ 333	\$ 2,198	\$ 12,105	\$ 19	(140,770)	\$ (8,601)
Net income	—	—	695	—	—	—
Cash dividends:						
Common (\$2.40 per share)	—	—	(485)	—	—	—
Common stock issued for:						
Share-based compensation and other plans, net	—	30	(2)	1	2,839	(6)
Acquisitions	—	2,083	—	—	15,959	368
Share-based compensation	—	164	—	—	—	—
Common stock held in trusts, net	—	—	—	—	5	—
Balance at September 30, 2015	\$ 333	\$ 4,475	\$ 12,314	\$ 20	(121,967)	\$ (8,239)
Net income	—	—	976	—	—	—
Cash dividends:						
Common (\$2.64 per share)	—	—	(562)	—	—	—
Common stock issued for:						
Share-based compensation and other plans, net	—	27	(1)	2	2,607	26
Share-based compensation	—	191	—	—	—	—
Common stock held in trusts, net	—	—	—	—	(11)	—
Balance at September 30, 2016	\$ 333	\$ 4,693	\$ 12,727	\$ 22	(119,371)	\$ (8,212)
Net income	—	—	1,100	—	—	—
Cash dividends:						
Common (\$2.92 per share)	—	—	(645)	—	—	—
Preferred	—	—	(70)	—	—	—
Common stock issued for:						
Public equity offerings	14	4,810	—	—	—	—
Share-based compensation and other plans, net	—	(65)	(1)	(3)	1,908	6
Share-based compensation	—	180	—	—	—	—
Common stock held in trusts, net	—	—	—	—	7	—
Repurchase of common stock	—	—	—	—	(1,289)	(220)
Balance at September 30, 2017	\$ 347	\$ 9,619	\$ 13,111	\$ 19	(118,745)	\$ (8,427)

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

Accelerated Share Repurchase Agreement

Using proceeds received from the divestiture of the Respiratory Solutions business in the first quarter of fiscal year 2017, the Company repurchased approximately 1.3 million shares of its common stock under an

accelerated share repurchase agreement. The repurchased shares were recorded as a \$220 million increase to *Common stock in treasury*.

Common and Preferred Stock Offerings

In May 2017 and in connection with the Company's pending agreement to acquire C.R. Bard, Inc. ("Bard"), which is further discussed in Note 9, the Company completed registered public offerings of equity securities including:

- 14.025 million shares of the Company's common stock for net proceeds of \$2.4 billion (gross proceeds of \$2.5 billion).
- 2.475 million shares of the Company's mandatory convertible preferred stock (ownership is held in the form of depositary shares, each representing a 1/20th interest in a share of preferred stock) for net proceeds of \$2.4 billion (gross proceeds of \$2.5 billion). If and when declared, dividends on the mandatory convertible preferred stock will be payable on a cumulative basis at an annual rate of 6.125% on the liquidation preference of \$1,000 per preferred share (\$50 per depositary share). The shares of preferred stock are convertible to a minimum of 11.7 million and up to a maximum of 14.0 million shares of Company common stock at an exchange ratio that is based on the market price of the Company's common stock at the date of conversion, and no later than the mandatory conversion date of May 1, 2020.

The Company will use the net proceeds from these offerings to finance a portion of the cash consideration payable upon the closing of the Bard acquisition, which the Company expects to occur in the fourth calendar quarter of 2017.

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

(Millions of dollars)	Total	Foreign Currency Translation	Benefit Plans	Cash Flow Hedges
Balance at September 30, 2014	\$ (1,001)	\$ (270)	\$ (705)	\$ (26)
Other comprehensive loss before reclassifications, net of taxes	(787)	(692)	(80)	(16)
Amounts reclassified into income, net of taxes	50	—	44	6
Balance at September 30, 2015	\$ (1,738)	\$ (961)	\$ (741)	\$ (36)
Other comprehensive loss before reclassifications, net of taxes	(251)	(50)	(190)	(11)
Amounts reclassified into income, net of taxes	60	—	48	12
Balance at September 30, 2016	\$ (1,929)	\$ (1,011)	\$ (883)	\$ (35)
Other comprehensive income before reclassifications, net of taxes	140	11	121	8
Amounts reclassified into income, net of taxes	66	—	58	8
Balance at September 30, 2017	<u>\$ (1,723)</u>	<u>\$ (1,001)</u>	<u>\$ (703)</u>	<u>\$ (18)</u>

The amount of foreign currency translation recognized in other comprehensive income during the year ended September 30, 2017 included net losses relating to net investment hedges, as further discussed in Note 13. The amount recognized in other comprehensive income during the year ended September 30, 2017 relating to cash flow hedges represented a net gain on forward starting interest rate swaps, which is further discussed in Note 13.

The tax impacts for amounts recognized in other comprehensive income before reclassifications were as follows:

(Millions of dollars)	<u>2017</u>	<u>2016</u>	<u>2015</u>
<i>Benefit Plans</i>			
Income tax (provision) benefit for net gains (losses) recorded in other comprehensive income	\$ (60)	\$ 79	\$ 47
<i>Cash Flow Hedges</i>			
Income tax (provision) benefit for net gains (losses) recorded in other comprehensive income	\$ (5)	\$ 7	\$ 10

Reclassifications out of *Accumulated other comprehensive income (loss)* relating to benefit plans and cash flow hedges in 2017, 2016 and 2015 were immaterial to the Company's consolidated financial results

Note 4 — Earnings per Share

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

	<u>2017</u>	<u>2016</u>	<u>2015</u>
Average common shares outstanding	218,943	212,702	202,537
Dilutive share equivalents from share-based plans (A) (B)	4,645	4,834	4,972
Average common and common equivalent shares outstanding — assuming dilution	<u>223,588</u>	<u>217,536</u>	<u>207,509</u>

- (A) For the year ended September 30, 2017, 5 million dilutive share equivalents associated with mandatory convertible preferred stock issued during 2017, as further discussed in Note 3, were excluded from the diluted shares outstanding calculation because the result would have been antidilutive. For the years ended September 30, 2017, 2016 and 2015 there were no options to purchase shares of common stock which were excluded from the diluted earnings per share calculation.
- (B) The adjustments to calculate diluted share equivalents from share-based plans in 2016 and 2015 included excess tax benefits relating to share-based compensation awards. Upon the Company's adoption, as discussed in Note 2, of new accounting requirements relating to share-based compensation award-related income tax effects, the adjustment in 2017 excluded these excess tax benefits.

Note 5 — Commitments and Contingencies

Commitments

Rental expense for all operating leases amounted to \$110 million in 2017, \$112 million in 2016 and \$89 million in 2015. Future minimum rental commitments on non-cancelable leases are as follows: 2018 — \$67 million; 2019 — \$57 million; 2020 — \$47 million; 2021 — \$35 million; 2022 — \$31 million and an aggregate of \$39 million thereafter.

As of September 30, 2017, the Company has certain future purchase commitments aggregating to approximately \$1.077 billion, which will be expended over the next several years.

Contingencies

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

In June 2007, Retractable Technologies, Inc. ("RTI") filed a complaint against the Company under the caption Retractable Technologies, Inc. vs. Becton Dickinson and Company (Civil Action No. 2:07-cv-250, U.S. District Court, Eastern District of Texas) alleging that the BD Integra™ syringes infringe patents licensed exclusively to RTI. In its complaint, RTI also alleged that the Company engaged in false advertising with respect to certain of the Company's safety-engineered products in violation of the Lanham Act; acted to exclude RTI from various product markets and to maintain its market share through, among other things, exclusionary contracts in violation of state and federal antitrust laws; and engaged in unfair competition. In January 2008, the Court severed the patent and non-patent claims into separate cases, and stayed the non-patent claims during the pendency of the patent claims at the trial court level. On April 1, 2008, RTI filed a complaint against BD under the caption Retractable Technologies, Inc. and Thomas J. Shaw v. Becton Dickinson and Company (Civil Action No. 2:08-cv-141, U.S. District Court, Eastern District of Texas) alleging that the BD Integra™ syringes infringe another patent licensed exclusively to RTI. On August 29, 2008, the Court ordered the consolidation of the patent cases. RTI was subsequently awarded \$5 million in damages at a jury trial with respect to the patent claims, which has been paid, and the patent cases are now concluded.

On September 19, 2013, a jury returned a verdict against BD with respect to RTI's Lanham Act claim and claim for attempted monopolization based on deception in the safety syringe market. The jury awarded RTI \$113.5 million for its attempted monopolization claim (which would be trebled under the antitrust statute). The jury's verdict rejected RTI's monopolization claims in the markets for safety syringes, conventional syringes and safety IV catheters; its attempted monopolization claims in the markets for conventional syringes and safety IV catheters; and its claims for contractual restraint of trade and exclusive dealing in the markets for safety syringes, conventional syringes and safety IV catheters. In connection with the verdict, the Company recorded a pre-tax charge of approximately \$341 million in the fourth quarter of fiscal year 2013. With respect to RTI's requested injunction relief, in November 2014, the Court granted RTI's request that BD be ordered to issue certain corrective statements regarding its advertising and enjoined from making certain advertising claims. The Court denied RTI's request for injunctive relief relating to BD's contracting practices and BD's safety syringe advertising, finding that RTI failed to prove that BD's contracting practices violated the antitrust laws or that BD's safety syringe advertising is false. On January 14, 2015, the Court granted in part and denied in part BD's motion for a stay of the injunction. The Court held that, pending appeal, BD would not be required to send the corrective advertising notices to end-user customers, but only to employees, distributors and Group Purchasing Organizations. On January 15, 2015, the Court entered its Final Judgment in the case ordering that RTI recover \$341 million for its attempted monopolization claim and \$12 million for attorneys' fees, and awarded pre and post-judgment interest and costs. On February 3, 2015, the Court of Appeals for the Fifth Circuit denied BD's motion for a stay of the injunction pending the final appeal, and BD thereafter complied with the Court's order. On April 23, 2015, the Court granted BD's motion to eliminate the award of pre-judgment interest, and entered a new Final Judgment. BD thereafter appealed to the Court of Appeals challenging the entirety of the Final Judgment. On December 2, 2016, the Court of Appeals issued an opinion reversing the judgment as to RTI's attempted monopolization claim and rendered judgment on that claim in favor of BD. As a result, the Company reversed \$336 million of reserves associated with this judgment, which was recorded in *Other operating (income) expense, net*. The Court of Appeals affirmed the judgment for Lanham Act liability, and remanded the case to the district court to consider whether and if so how much profit should be disgorged by BD on that claim. The Court of Appeals vacated and remanded the injunction ordered by the Court. On January 31, 2017, RTI filed a petition for a writ of certiorari with the U.S. Supreme Court.

On March 20, 2017, the U.S. Supreme Court denied certiorari, and the district court thereafter heard RTI's request for disgorgement. On August 17, 2017, the district court entered judgment in favor of BD and ruled that RTI is not entitled to any award of money damages. RTI has appealed this ruling to the Fifth Circuit Court of Appeals.

On July 17, 2015, a class action complaint was filed against the Company in the U.S. District Court for the Southern District of Georgia. The plaintiffs, Glynn-Brunswick Hospital Authority, trading as Southeast Georgia Health System, and Southeast Georgia Health System, Inc., seek to represent a class of acute care purchasers of BD syringes and IV catheters. The complaint alleges that BD monopolized the markets for syringes and IV catheters through contracts, theft of technology, false advertising, acquisitions, and other conduct. The complaint seeks treble damages but does not specify the amount of alleged damages. The Company filed a motion to dismiss the complaint which was granted on January 29, 2016. On September 23, 2016, the court denied plaintiffs' motion to alter or amend the judgment to allow plaintiffs to file an amended complaint, and plaintiffs appealed that decision to the Eleventh Circuit Court of Appeals. The plaintiffs thereafter voluntarily dismissed their appeal, and the Court of Appeals dismissed the case on November 21, 2016.

The Company is also involved both as a plaintiff and a defendant in other legal proceedings and claims that arise in the ordinary course of business. The Company believes that it has meritorious defenses to the suits pending against the Company and is engaged in a vigorous defense of each of these matters.

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

Note 6 — Segment Data

The Company’s organizational structure is based upon two principal business segments: BD Medical (“Medical”) and BD Life Sciences (“Life Sciences”). The Company’s two principal business segments are strategic businesses that are managed separately because each one develops, manufactures and markets distinct products and services.

BD Medical

BD Medical produces a broad array of medical technologies and devices that are used to help improve healthcare delivery in a wide range of settings. The primary customers served by BD Medical are hospitals and clinics; physicians’ office practices; consumers and retail pharmacies; governmental and nonprofit public health agencies; pharmaceutical companies; and healthcare workers. BD Medical consists of the following organizational units:

Organizational Unit	Principal Product Lines
Diabetes Care	Syringes, pen needles and other products related to the injection or infusion of insulin and other drugs used in the treatment of diabetes.
Medication and Procedural Solutions	Needles, syringes and intravenous catheters for medication delivery (including safety-engineered and auto-disable devices); prefilled IV flush syringes; regional anesthesia needles and trays; sharps disposal containers; closed-system transfer devices; skin antiseptic products; and surgical and laproscopic instrumentation.
Medication Management Solutions	Intravenous medication safety and infusion therapy delivery systems, including infusion pumps and dedicated disposables; medication compounding workflow systems; and automated medication dispensing; automated supply management systems; medication inventory optimization and tracking systems; and analytics related to all the above products.
Pharmaceutical Systems	Prefillable drug delivery systems provided to pharmaceutical companies for use as containers for injectable pharmaceutical products, which are then placed on the market as drug/device combinations.

BD Life Sciences

BD Life Sciences provides products for the safe collection and transport of diagnostics specimens, and instruments and reagent systems to detect a broad range of infectious diseases, healthcare-associated infections (“HAIs”) and cancers. In addition, BD Life Sciences produces research and clinical tools that facilitate the study of cells, and the components of cells, to gain a better understanding of normal and disease processes. That information is used to aid the discovery and development of new drugs and vaccines, and to improve the diagnosis and management of diseases. The primary customers served by BD Life Sciences are hospitals, laboratories and clinics; blood banks; healthcare workers; public health agencies; physicians’ office practices; academic and government institutions; and pharmaceutical and biotechnology companies. BD Life Sciences consists of the following organizational units:

Organizational Unit	Principal Product Lines
Preanalytical Systems	Integrated systems for specimen collection; safety-engineered blood collection products and systems.
Diagnostic Systems	Automated blood culturing and tuberculosis culturing systems; molecular testing systems for infectious diseases and women's health; microorganism identification and drug susceptibility systems; liquid-based cytology systems for cervical cancer screening; rapid diagnostic assays; microbiology laboratory automation; and plated media.
Biosciences	Fluorescence-activated cell sorters and analyzers; monoclonal antibodies and kits for performing cell analysis; reagent systems for life science research; molecular indexing and next-generation sequencing sample preparation for genomics research; clinical oncology, immunological (HIV) and transplantation diagnostic/monitoring reagents and analyzers; and cell culture media supplements for biopharmaceutical manufacturing.

Distribution of products is primarily through independent distribution channels, and directly to end-users by BD and independent sales representatives. No customer accounted for 10% or more of revenues in any of the three years presented.

The Company evaluates performance of its business segments and allocates resources to them primarily based upon operating income. Segment operating income represents revenues reduced by product costs and operating expenses. As more fully discussed in Note 10, the Company sold a 50.1% controlling financial interest in its Respiratory Solutions business, a component of the Medical segment, in October 2016. This transaction did not meet the criteria established for reporting discontinued operations and as such, results for the years ended September 30, 2016 and 2015 included \$822 million and \$417 million, respectively, of revenues which did not occur in the current year.

(Millions of dollars)	2017	2016	2015
Revenues (A)			
Medical	\$ 8,105	\$ 8,654	\$ 6,460
Life Sciences	3,988	3,829	3,822
Total Revenues	<u>\$ 12,093</u>	<u>\$ 12,483</u>	<u>\$ 10,282</u>
Income Before Income Taxes			
Medical	\$ 2,155	\$ 2,052	\$ 1,530
Life Sciences	772	793	839
Total Segment Operating Income	<u>2,927</u>	<u>2,845</u>	<u>2,368</u>
Acquisitions and other restructurings	(354)	(728)	(426)
Net interest expense	(445)	(367)	(356)
Other unallocated items (B)	(1,152)	(676)	(847)
Income Before Income Taxes	<u>\$ 976</u>	<u>\$ 1,074</u>	<u>\$ 739</u>
Segment Assets			
Medical	\$ 18,332	\$ 19,154	\$ 20,055
Life Sciences	4,056	3,848	3,932
Total Segment Assets	<u>22,388</u>	<u>23,002</u>	<u>23,987</u>
Corporate and All Other (C)	<u>15,347</u>	<u>2,584</u>	<u>2,491</u>
Total Assets	<u>\$ 37,734</u>	<u>\$ 25,586</u>	<u>\$ 26,478</u>
Capital Expenditures			
Medical	\$ 502	\$ 482	\$ 414
Life Sciences	212	200	168
Corporate and All Other	13	12	14
Total Capital Expenditures	<u>\$ 727</u>	<u>\$ 693</u>	<u>\$ 596</u>
Depreciation and Amortization			
Medical	\$ 825	\$ 857	\$ 619
Life Sciences	254	254	256
Corporate and All Other	10	3	17
Total Depreciation and Amortization	<u>\$ 1,088</u>	<u>\$ 1,114</u>	<u>\$ 891</u>

(A) Intersegment revenues are not material.

(B) Primarily comprised of foreign exchange, corporate expenses, and share-based compensation expense. Results in 2017 included a \$748 million non-cash charge resulting from a modification to the Company's dispensing equipment lease contracts with customers, as well as a \$336 million reversal of certain reserves related to an appellate court decision which, among other things, reversed an unfavorable antitrust judgment in the RTI case. Additional disclosures regarding the legal matter and the lease contract modification are provided in Notes 5 and 17, respectively. Results in 2015 reflected \$293 million in recognition of the fair value step-up adjustment recorded relative to CareFusion's inventory on the acquisition date.

(C) Includes cash and investments and corporate assets.

Geographic Information

The countries in which the Company has local revenue-generating operations have been combined into the following geographic areas: the United States (including Puerto Rico); Europe; Greater Asia (which includes

Japan and Asia Pacific); and Other, which is comprised of Latin America, Canada, and EMA (which includes the Commonwealth of Independent States, Middle East and Africa).

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

(Millions of dollars)	2017	2016	2015
Revenues			
United States	\$ 6,504	\$ 6,893	\$ 5,069
Europe	2,588	2,674	2,434
Greater Asia	1,744	1,692	1,545
Other	1,257	1,225	1,234
	<u>\$ 12,093</u>	<u>\$ 12,483</u>	<u>\$ 10,282</u>
Long-Lived Assets			
United States	\$ 13,151	\$ 14,075	\$ 15,513
Europe	4,421	3,747	3,908
Greater Asia	578	586	573
Other	584	483	494
Corporate	366	329	332
	<u>\$ 19,101</u>	<u>\$ 19,220</u>	<u>\$ 20,819</u>

Note 7 — Share-Based Compensation

The Company grants share-based awards under the 2004 Employee and Director Equity-Based Compensation Plan (“2004 Plan”), which provides long-term incentive compensation to employees and directors consisting of: stock appreciation rights (“SARs”), stock options, performance-based restricted stock units, time-vested restricted stock units and other stock awards.

The fair value of share-based payments is recognized as compensation expense in net income. The amounts and location of compensation cost relating to share-based payments included in the consolidated statements of income is as follows:

(Millions of dollars)	2017	2016	2015
Cost of products sold	\$ 30	\$ 29	\$ 23
Selling and administrative expense	113	106	82
Research and development expense	24	22	17
Acquisitions and other restructurings	10	39	44
	<u>\$ 177</u>	<u>\$ 196</u>	<u>\$ 166</u>
Tax benefit associated with share-based compensation costs recognized	<u>\$ 61</u>	<u>\$ 69</u>	<u>\$ 59</u>

In 2015, certain pre-acquisition equity awards of CareFusion were converted into BD restricted stock awards or BD stock options with accelerated vesting terms at the acquisition date. In addition, as an incentive to encourage post-acquisition employee retention, certain pre-acquisition equity awards of CareFusion were converted into either BD restricted stock awards or BD stock options, as applicable, as of the acquisition date, with substantially the same terms and conditions as were applicable under such CareFusion awards immediately prior to the acquisition date. The compensation expense associated with these replacement awards was recorded in *Acquisitions and other restructurings*.

Stock Appreciation Rights

SARs represent the right to receive, upon exercise, shares of common stock having a value equal to the difference between the market price of common stock on the date of exercise and the exercise price on the date of grant. SARs vest over a four-year period and have a ten-year term. The fair value was estimated on the date of grant using a lattice-based binomial option valuation model that uses the following weighted-average assumptions:

	<u>2017</u>	<u>2016</u>	<u>2015</u>
Risk-free interest rate	2.33%	2.17%	2.20%
Expected volatility	20.0%	19.0%	19.0%
Expected dividend yield	1.71%	1.76%	1.78%
Expected life	7.5 years	7.6 years	7.6 years
Fair value derived	\$33.81	\$27.69	\$24.82

Expected volatility is based upon historical volatility for the Company's common stock and other factors. The expected life of SARs granted is derived from the output of the lattice-based model, using assumed exercise rates based on historical exercise and termination patterns, and represents the period of time that SARs granted are expected to be outstanding. The risk-free interest rate used is based upon the published U.S. Treasury yield curve in effect at the time of grant for instruments with a similar life. The dividend yield is based upon the most recently declared quarterly dividend as of the grant date. The Company issued 793 thousand shares during 2017 to satisfy the SARs exercised.

A summary of SARs outstanding as of September 30, 2017 and changes during the year then ended is as follows:

	<u>SARs (in thousands)</u>	<u>Weighted Average Exercise Price</u>	<u>Weighted Average Remaining Contractual Term (Years)</u>	<u>Aggregate Intrinsic Value (Millions of dollars)</u>
Balance at October 1	7,027	\$ 103.83		
Granted	996	170.69		
Exercised	(1,445)	83.81		
Forfeited, canceled or expired	(112)	142.04		
Balance at September 30	<u>6,466</u>	<u>\$ 117.94</u>	<u>6.24</u>	<u>\$ 504</u>
Vested and expected to vest at September 30	<u>6,215</u>	<u>\$ 116.54</u>	<u>6.16</u>	<u>\$ 493</u>
Exercisable at September 30	<u>3,952</u>	<u>\$ 96.00</u>	<u>4.98</u>	<u>\$ 395</u>

A summary of SARs exercised 2017, 2016 and 2015 is as follows:

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Total intrinsic value of SARs exercised	<u>\$ 148</u>	<u>\$ 148</u>	<u>\$ 96</u>
Tax benefit realized from SAR exercises	<u>\$ 53</u>	<u>\$ 52</u>	<u>\$ 34</u>
Total fair value of SARs vested	<u>\$ 30</u>	<u>\$ 24</u>	<u>\$ 22</u>

Stock Options

The Company has not granted stock options since 2005. As previously discussed, certain pre-acquisition equity awards of CareFusion were converted on March 17, 2015 into BD stock options with accelerated vesting terms. A summary of these stock options outstanding as of September 30, 2017 and changes during the year then ended is as follows:

	Stock Options (in thousands)	Weighted Average Exercise Price	Weighted Average Remaining Contractual Term (Years)	Aggregate Intrinsic Value (Millions of dollars)
Balance at October 1	495	\$ 79.99		
Exercised	(186)	77.66		
Forfeited, canceled or expired	(6)	81.30		
Balance at September 30	303	\$ 81.40	2.56	\$ 35
Vested at September 30	303	\$ 81.40	2.56	\$ 35
Exercisable at September 30	303	\$ 81.40	2.56	\$ 35

A summary of options exercised 2017, 2016 and 2015 is as follows:

(Millions of dollars)	2017	2016	2015
Cash received from stock option exercises	\$ 14	\$ 50	\$ 75
Tax benefit realized from stock option exercises	\$ 8	\$ 17	\$ 20
Total intrinsic value of stock options exercised	\$ 20	\$ 51	\$ 52
Total fair value of stock options vested	\$ 2	\$ 11	\$ 59

Performance-Based and Time-Vested Restricted Stock Units

Performance-based restricted stock units cliff vest three years after the date of grant. These units are tied to the Company's performance against pre-established targets over a three-year performance period. The performance measures for fiscal years 2017, 2016 and 2015 were relative total shareholder return (measures the Company's stock performance during the performance period against that of peer companies) and average annual return on invested capital. Under the Company's long-term incentive program, the actual payout under these awards may vary from zero to 200% of an employee's target payout, based on the Company's actual performance over the three-year performance period. The fair value is based on the market price of the Company's stock on the date of grant. Compensation cost initially recognized assumes that the target payout level will be achieved and is adjusted for subsequent changes in the expected outcome of performance-related conditions. For units for which the performance conditions are modified after the date of grant, any incremental increase in the fair value of the modified units, over the original units, is recorded as compensation expense on the date of the modification for vested units, or over the remaining performance period for units not yet vested.

Time-vested restricted stock unit awards granted after January 2015 vest on a graded basis over a three-year period. Time-vested restricted stock units granted before January 2015 cliff vest three years after the date of grant, except for certain key executives of the Company, including the executive officers, for which such units generally vest one year following the employee's retirement. The related share-based compensation expense is recorded over the requisite service period, which is the vesting period or is based on retirement eligibility. The fair value of all time-vested restricted stock units is based on the market value of the Company's stock on the date of grant.

A summary of restricted stock units outstanding as of September 30, 2017 and changes during the year then ended is as follows:

	Performance-Based		Time-Vested	
	Stock Units (in thousands)	Weighted Average Grant Date Fair Value	Stock Units (in thousands)	Weighted Average Grant Date Fair Value
Balance at October 1	1,112	\$ 148.27	2,584	\$ 123.16
Granted	381	174.92	845	165.96
Distributed	(145)	134.19	(799)	117.06
Forfeited or canceled	(268)	139.93	(494)	124.52
Balance at September 30	1,080 (A)	\$ 161.64	2,136	\$ 142.06
Expected to vest at September 30	428 (B)	\$ 159.22	2,052	\$ 141.36

(A) Based on 200% of target payout.

(B) Net of expected forfeited units and units in excess of the expected performance payout of 79 thousand and 573 thousand shares, respectively.

The weighted average grant date fair value of restricted stock units granted during the years 2017, 2016 and 2015 are as follows:

	Performance-Based			Time-Vested		
	2017	2016	2015	2017	2016	2015
Weighted average grant date fair value of units granted	\$174.92	\$153.73	\$156.65	\$165.96	\$145.57	\$136.30

The total fair value of stock units vested during 2017, 2016 and 2015 was as follows:

(Millions of dollars)	Performance-Based			Time-Vested		
	2017	2016	2015	2017	2016	2015
Total fair value of units vested	\$ 32	\$ 22	\$ 16	\$ 139	\$ 114	\$ 181

At September 30, 2017, the weighted average remaining vesting term of performance-based and time vested restricted stock units is 1.22 and 1 year, respectively.

Unrecognized Compensation Expense and Other Stock Plans

The amount of unrecognized compensation expense for all non-vested share-based awards as of September 30, 2017, is approximately \$182 million, which is expected to be recognized over a weighted-average remaining life of approximately 1.93 years. At September 30, 2017, 8.8 million shares were authorized for future grants under the 2004 Plan. The Company has a policy of satisfying share-based payments through either open market purchases or shares held in treasury. At September 30, 2017, the Company has sufficient shares held in treasury to satisfy these payments.

At September 30, 2017 and 2016, awards for 29 thousand and 43 thousand shares, respectively, were outstanding under a plan, that was terminated in 2004, which allowed for grants of common shares to certain key employees. The Company has a Directors' Deferral Plan, which provides a means to defer director compensation, from time to time, on a deferred stock or cash basis. As of September 30, 2017, 130 thousand shares were held in trust, of which one thousand shares represented Directors' compensation in 2017, in accordance with the provisions of the plan. Under this plan, which is unfunded, directors have an unsecured contractual commitment from the Company. The Company also has a Deferred Compensation Plan that allows

certain highly-compensated employees, including executive officers, to defer salary, annual incentive awards and certain equity-based compensation. As of September 30, 2017, 338 thousand shares were issuable under this plan.

Note 8 — Benefit Plans

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

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

(Millions of dollars)	Pension Plans			Other Postretirement Benefits		
	2017	2016	2015	2017	2016	2015
Service cost	\$ 110	\$ 81	\$ 77	\$ 3	\$ 3	\$ 3
Interest cost	61	72	87	4	5	7
Expected return on plan assets	(112)	(109)	(123)	—	—	—
Amortization of prior service credit	(14)	(15)	(15)	(5)	(5)	(5)
Amortization of loss	92	77	68	2	2	3
Settlements	—	7	—	—	—	—
Net pension and postretirement cost	<u>\$ 138</u>	<u>\$ 113</u>	<u>\$ 93</u>	<u>\$ 4</u>	<u>\$ 5</u>	<u>\$ 9</u>
Net pension cost attributable to international plans	<u>\$ 43</u>	<u>\$ 35</u>	<u>\$ 32</u>			

The amounts provided above for amortization of prior service credit and amortization of loss represent the reclassifications of prior service credits and net actuarial losses that were recognized in *Accumulated other comprehensive income (loss)* in prior periods. The settlement loss recorded in 2016 primarily included lump sum benefit payments associated with the Company's U.S. supplemental pension plan. The Company recognizes pension settlements when payments from the supplemental plan exceed the sum of service and interest cost components of net periodic pension cost associated with this plan for the fiscal year.

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

(Millions of dollars)	Pension Plans		Other Postretirement Benefits	
	2017	2016	2017	2016
Change in benefit obligation:				
Beginning obligation	\$ 2,719	\$ 2,426	\$ 184	\$ 186
Service cost	110	81	3	3
Interest cost	61	72	4	5
Plan amendments	(1)	—	—	(1)
Benefits paid	(123)	(116)	(16)	(17)
Impact of divestitures	(19)	—	—	—
Actuarial (gain) loss	(134)	302	(15)	3
Settlements	(1)	(15)	—	—
Other, includes translation	36	(30)	4	4
Benefit obligation at September 30	<u>\$ 2,647</u>	<u>\$ 2,719</u>	<u>\$ 165</u>	<u>\$ 184</u>
Change in fair value of plan assets:				
Beginning fair value	\$ 1,855	\$ 1,732	\$ —	\$ —
Actual return on plan assets	134	131	—	—
Employer contribution	54	145	—	—
Benefits paid	(123)	(116)	—	—
Impact of divestitures	(13)	—	—	—
Settlements	(1)	(15)	—	—
Other, includes translation	26	(21)	—	—
Plan assets at September 30	<u>\$ 1,932</u>	<u>\$ 1,855</u>	<u>\$ —</u>	<u>\$ —</u>
Funded Status at September 30:				
Unfunded benefit obligation	<u>\$ (715)</u>	<u>\$ (864)</u>	<u>\$ (165)</u>	<u>\$ (184)</u>
Amounts recognized in the Consolidated Balance Sheets at September 30:				
Other	\$ 9	\$ 5	\$ —	\$ —
Salaries, wages and related items	(17)	(12)	(14)	(15)
Long-term Employee Benefit Obligations	(707)	(857)	(150)	(169)
Net amount recognized	<u>\$ (715)</u>	<u>\$ (864)</u>	<u>\$ (165)</u>	<u>\$ (184)</u>
Amounts recognized in Accumulated other comprehensive income (loss) before income taxes at September 30:				
Prior service credit	74	87	28	33
Net actuarial loss	(1,065)	(1,307)	(15)	(32)
Net amount recognized	<u>\$ (991)</u>	<u>\$ (1,221)</u>	<u>\$ 14</u>	<u>\$ 1</u>

International pension plan assets at fair value included in the preceding table were \$678 million and \$624 million at September 30, 2017 and 2016, respectively. The international pension plan projected benefit obligations were \$917 million and \$951 million at September 30, 2017 and 2016, respectively.

Pension plans with accumulated benefit obligations in excess of plan assets and plans with projected benefit obligations in excess of plan assets consist of the following at September 30:

(Millions of dollars)	Accumulated Benefit Obligation Exceeds the Fair Value of Plan Assets		Projected Benefit Obligation Exceeds the Fair Value of Plan Assets	
	2017	2016	2017	2016
Projected benefit obligation	\$ 2,551	\$ 2,616	\$ 2,613	\$ 2,682
Accumulated benefit obligation	\$ 2,470	\$ 2,529		
Fair value of plan assets	\$ 1,833	\$ 1,757	\$ 1,889	\$ 1,813

The estimated net actuarial loss and prior service credit for pension benefits that will be amortized from *Accumulated other comprehensive income (loss)* into net pension costs over the next fiscal year are expected to be \$76 million and \$13 million, respectively. The net actuarial loss for other postretirement benefits that will be amortized from *Accumulated other comprehensive income (loss)* into net other postretirement costs over the next fiscal year is immaterial. The estimated prior service credit that will be amortized from *Accumulated other comprehensive income (loss)* into net other postretirement costs over the next fiscal year is expected to be \$5 million.

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

	2017	2016	2015
Net Cost			
Discount rate:			
U.S. plans (A)	3.42%	4.15%	4.15%
International plans	1.70	2.84	3.14
Expected return on plan assets:			
U.S. plans	7.25	7.50	7.50
International plans	4.65	5.02	5.45
Rate of compensation increase:			
U.S. plans	4.25	4.25	4.25
International plans	2.33	2.33	2.49
Benefit Obligation			
Discount rate:			
U.S. plans	3.72	3.42	4.15
International plans	2.25	1.70	2.84
Rate of compensation increase:			
U.S. plans	4.51	4.25	4.25
International plans	2.30	2.33	2.33

(A) In 2015 the Company calculated the service and interest components utilizing a single weighted-average discount rate derived from the yield curve used to measure the benefit obligation at the beginning of the period. Effective September 30, 2015, the Company elected to utilize an approach that discounts the individual expected cash flows using the applicable spot rates derived from the yield curve over the projected cash flow period. The Company accounted for this change as a change in accounting estimate that is inseparable from a change in accounting principle and as such, the change was accounted for prospectively.

At September 30, 2017 the assumed healthcare trend rates were 7.0%, gradually decreasing to an ultimate rate of 5.0% beginning in 2027. At September 30, 2016 the assumed healthcare trend rates were 6.6% pre and post age 65, gradually decreasing to an ultimate rate of 5.0% beginning in 2024. A one percentage point

increase or decrease in assumed healthcare cost trend rates in each year would not materially impact the accumulated postretirement benefit obligation as of September 30, 2017 or the aggregate of the service cost and interest cost components of 2017 annual expense.

Expected Rate of Return on Plan Assets

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

Expected Funding

The Company's funding policy for its defined benefit pension plans is to contribute amounts sufficient to meet legal funding requirements, plus any additional amounts that may be appropriate considering the funded status of the plans, tax consequences, the cash flow generated by the Company and other factors. The Company made a discretionary contribution of \$112 million to its U.S. pension plan in October 2017. The Company does not anticipate any significant required contributions to its pension plans in 2018.

Expected benefit payments are as follows:

(Millions of dollars)	Pension Plans	Other Postretirement Benefits
2018	\$ 173	\$ 14
2019	158	14
2020	159	14
2021	163	13
2022	165	13
2023-2027	859	56

Investments

The Company's primary objective is to achieve returns sufficient to meet future benefit obligations. It seeks to generate above market returns by investing in more volatile asset classes such as equities while at the same time controlling risk through diversification in non-correlated asset classes and through allocations to more stable asset classes like fixed income.

U.S. Plans

The Company's U.S. pension plans comprise 65% of total benefit plan investments, based on September 30, 2017 market values and have a target asset mix of 40% fixed income, 28% diversifying investments and 32% equities. This mix was established based on an analysis of projected benefit payments and estimates of long-term returns, volatilities and correlations for various asset classes. The asset allocations to diversifying investments include high-yield bonds, hedge funds, real estate, infrastructure, commodities, leveraged loans and emerging markets bonds.

The actual portfolio investment mix may, from time to time, deviate from the established target mix due to various factors such as normal market fluctuations, the reliance on estimates in connection with the determination of allocations and normal portfolio activity such as additions and withdrawals. Rebalancing of the asset portfolio on a quarterly basis is required to address any allocations that deviate from the established target allocations in excess of defined allowable ranges. The target allocations are subject to periodic review, including a review of the asset portfolio's performance, by the named fiduciary of the plans. Any tactical deviations from the established asset mix require the approval of the named fiduciary.

The U.S. plans may enter into both exchange traded and non-exchange traded derivative transactions in order to manage interest rate exposure, volatility, term structure of interest rates, and sector and currency

exposures within the fixed income portfolios. The Company has established minimum credit quality standards for counterparties in such transactions.

The following table provides the fair value measurements of U.S. plan assets, as well as the measurement techniques and inputs utilized to measure fair value of these assets, at September 30, 2017 and 2016. The categorization of fund investments is based upon the categorization of these funds' underlying assets.

(Millions of dollars)	Total U.S. Plan Asset Balances		Investments Measured at Net Asset Value (A)		Quoted Prices in Active Markets for Identical Assets (Level 1)		Significant Other Observable Inputs (Level 2)		Significant Unobservable Inputs (Level 3)	
	2017	2016	2017	2016	2017	2016	2017	2016	2017	2016
Fixed Income:										
Mortgage and asset-backed securities	\$ 155	\$ 169	\$ —	\$ —	\$ —	\$ —	\$ 155	\$ 169	\$ —	\$ —
Corporate bonds	232	197	—	—	89	68	144	129	—	—
Government and agency-U.S.	107	103	—	—	83	67	25	36	—	—
Government and agency-Foreign	98	90	12	32	63	52	22	6	—	—
Equity securities	369	348	307	287	62	61	—	—	—	—
Cash and cash equivalents	40	89	—	—	40	89	—	—	—	—
Other	252	234	217	201	34	33	—	—	—	—
Fair value of plan assets	<u>\$1,254</u>	<u>\$1,231</u>	<u>\$ 537</u>	<u>\$ 520</u>	<u>\$ 371</u>	<u>\$ 371</u>	<u>\$ 346</u>	<u>\$ 340</u>	<u>\$ —</u>	<u>\$ —</u>

(A) As per applicable disclosure requirements, certain investments that were measured at net asset value per share or its equivalent have not been categorized within the fair value hierarchy. Values of such assets are based on the corroborated net asset value provided by the fund administrator.

Fixed Income Securities

U.S. pension plan assets categorized above as fixed income securities include fund investments comprised of mortgage-backed, corporate, government and agency and asset-backed instruments. Mortgage-backed securities consist of residential mortgage pass-through certificates. Investments in corporate bonds are diversified across industry and sector and consist of investment-grade, as well as high-yield debt instruments. U.S. government investments consist of obligations of the U.S. Treasury, other U.S. government agencies, state governments and local municipalities. Assets categorized as foreign government and agency debt securities included investments in developed and emerging markets.

The values of fixed income investments classified within Level 1 are based on the closing price reported on the major market on which the investments are traded. A portion of the fixed income instruments classified within Level 2 are valued based upon estimated prices from independent vendors' pricing models and these prices are derived from market observable sources including: benchmark yields, reported trades, broker/dealer quotes, issuer spreads, benchmark securities, bids, offers and other market-related data.

Equity Securities

U.S. pension plan assets categorized as equity securities consist of fund investments in publicly-traded U.S. and non-U.S. equity securities. In order to achieve appropriate diversification, these portfolios are invested

across market sectors, investment styles, capitalization weights and geographic regions. The values of equity securities classified within Level 1 are based on the closing price reported on the major market on which the investments are traded.

Cash and Cash Equivalents

A portion of the U.S. plans' assets consists of investments in cash and cash equivalents, primarily to accommodate liquidity requirements relating to trade settlement and benefit payment activity, and the values of these assets are based upon quoted market prices.

Other Securities

Other U.S. pension plan assets include fund investments comprised of underlying assets of real estate, infrastructure, commodities and hedge funds. The values of such instruments classified within Level 1 are based on the closing price reported on the major market on which the investments are traded.

International Plans

International plan assets comprise 35% of the Company's total benefit plan assets, based on market value at September 30, 2017. Such plans have local independent fiduciary committees, with responsibility for development and oversight of investment policy, including asset allocation decisions. In making such decisions, consideration is given to local regulations, investment practices and funding rules.

The following table provides the fair value measurements of international plan assets, as well as the measurement techniques and inputs utilized to measure fair value of these assets, at September 30, 2017 and 2016.

(Millions of dollars)	Total International Plan Asset Balances		Investments Measured at Net Asset Value (A)		Quoted Prices in Active Markets for Identical Assets (Level 1)		Significant Other Observable Inputs (Level 2)		Significant Unobservable Inputs (Level 3)	
	2017	2016	2017	2016	2017	2016	2017	2016	2017	2016
Fixed Income:										
Corporate bonds	\$ 14	\$ 34	\$ —	\$ —	\$ —	\$ —	\$ 13	\$ 34	\$ —	\$ —
Government and agency-U.S.	5	5	—	—	1	2	3	3	—	—
Government and agency-Foreign	127	119	—	—	83	73	45	46	—	—
Other fixed income	64	51	—	—	57	46	7	5	—	—
Equity securities	256	228	13	14	242	214	—	—	—	—
Cash and cash equivalents	28	13	—	—	28	13	—	—	—	—
Real estate	26	17	—	—	—	—	26	17	—	—
Insurance contracts	98	102	—	—	—	—	—	—	98	102
Other	62	57	—	—	47	43	15	14	—	—
Fair value of plan assets	<u>\$ 678</u>	<u>\$ 624</u>	<u>\$ 13</u>	<u>\$ 14</u>	<u>\$ 459</u>	<u>\$ 391</u>	<u>\$ 108</u>	<u>\$ 119</u>	<u>\$ 98</u>	<u>\$ 102</u>

- (A) As per applicable disclosure requirements, certain investments that were measured at net asset value per share or its equivalent have not been categorized within the fair value hierarchy. Values of such assets are based on the corroborated net asset value provided by the fund administrator.

Fixed Income Securities

Fixed income investments held by international pension plans include corporate, U.S. government and non-U.S. government securities. The values of fixed income securities classified within Level 1 are based on the closing price reported on the major market on which the investments are traded. Values of investments classified within Level 2 are based upon estimated prices from independent vendors' pricing models and these prices are derived from market observable sources.

Equity Securities

Equity securities included in the international plan assets consist of publicly-traded U.S. and non-U.S. equity securities. The values of equity securities classified within Level 1 are based on the closing price reported on the major market on which the investments are traded.

Other Securities

The international plans hold a portion of assets in cash and cash equivalents, in order to accommodate liquidity requirements and the values are based upon quoted market prices. Real estate investments consist of investments in funds holding an interest in real properties and the corresponding values represent the estimated fair value based on the fair value of the underlying investment value or cost, adjusted for any accumulated earnings or losses. The values of insurance contracts approximately represent cash surrender value. Other investments include fund investments for which values are based upon either quoted market prices or market observable sources.

The following table summarizes the changes, for the years ended September 30, 2017 and 2016, in the fair value of international pension assets measured using Level 3 inputs:

(Millions of dollars)	<u>Insurance Contracts</u>
Balance at September 30, 2015	\$ 90
Actual return on plan assets:	
Relating to assets held at September 30, 2015	8
Purchases, sales and settlements, net	2
Exchange rate changes	1
Balance at September 30, 2016	<u>\$ 102</u>
Actual return on plan assets:	
Relating to assets held at September 30, 2016	1
Purchases, sales and settlements, net	(11)
Transfers in from other categories	1
Exchange rate changes	4
Balance at September 30, 2017	<u><u>\$ 98</u></u>

Savings Incentive Plan

The Company has voluntary defined contribution plans covering eligible employees in the United States which provide for a Company match. The cost of these plans was \$67 million in 2017, \$61 million in 2016 and \$54 million in 2015.

Note 9 – Acquisitions

Definitive Agreement to Acquire Bard

On April 23, 2017, the Company announced that it had entered into a definitive agreement under which BD will acquire Bard for an implied value of \$317.00 per Bard common share in cash and stock, for estimated total consideration of approximately \$24 billion. The combination will create a highly differentiated medical technology company uniquely positioned to improve both the process of care and the treatment of disease for patients and healthcare providers.

Under the terms of the transaction, Bard common shareholders will be entitled to receive approximately \$222.93 in cash and 0.5077 shares of BD common stock per Bard share, or an implied value of \$317.00 per Bard common share based on BD's closing price on April 21, 2017. At closing, Bard shareholders will own approximately 15 percent of the combined company. The Company will finance the cash portion of total consideration transferred with available cash, which will include \$4.8 billion of net proceeds raised in the third quarter of fiscal year 2017 through registered public offerings of equity securities and approximately \$9.6 billion of net proceeds also raised in the third quarter through debt transactions. The total consideration transferred will also include an estimated \$8 billion of BD common stock. The transaction is subject to regulatory approval and customary closing conditions, and is expected to close in the fourth calendar quarter of 2017.

CareFusion Corporation

Overview of Transaction and Consideration Transferred

On March 17, 2015, the Company acquired a 100% interest in CareFusion, a global medical technology company with a comprehensive portfolio of products in the areas of medication management, infection prevention, operating room and procedural effectiveness, and respiratory care, to create a global leader in medication management and patient safety solutions. The fair value of consideration transferred consisted of the components below.

(Millions of dollars)	
Cash consideration	\$ 10,085
Noncash consideration-fair value of shares issued	2,269
Noncash consideration-fair value of stock options and other equity awards	184
Total consideration transferred	<u>\$ 12,538</u>

Additional disclosures regarding the financing arrangements the Company entered into to fund the cash portion of the consideration transferred relative to this acquisition are provided in Note 15.

Unaudited Pro Forma Information

The acquisition was accounted for under the acquisition method of accounting for business combinations. The operating activities from the acquisition date through March 31, 2015 were not material to the Company's consolidated results of operations. As such, CareFusion's operating results were included in the Company's consolidated results of operations beginning on April 1, 2015. *Revenues* and *Operating Income* are no longer specifically identifiable due to the progression of the Company's integration activities. *Revenues* and *Operating Income* for the year ended September 30, 2015 included revenues and operating loss attributable to CareFusion of \$2 billion and \$242 million, respectively.

The following table provides the pro forma results for the years ended September 30, 2016 and 2015 as if CareFusion had been acquired as of October 1, 2013.

(Millions of dollars, except per share data)

	2016	2015
Revenues	\$ 12,497	\$ 12,368
Net Income	\$ 1,453	\$ 1,276
Diluted Earnings per Share	\$ 6.68	\$ 5.92

The pro forma net income amounts above reflect adjustments, which were adjusted for the applicable tax impact, including the following: additional amortization and depreciation expense relating to assets acquired; interest and other financing costs relating to the acquisition transaction; and the elimination of one-time or nonrecurring items. The pro forma results do not include any anticipated cost savings or other effects of the integration of CareFusion. Accordingly, the pro forma results above are not necessarily indicative of the results that would have been if the acquisition had occurred on the dates indicated.

Other Transactions

Fiscal year 2017 acquisition of remaining interest in Caesarea Medical Electronics

Upon the Company's acquisition of CareFusion, it acquired a 40% ownership interest in Caesarea Medical Electronics ("CME"), an Israeli-based global infusion pump systems manufacturer. The Company previously accounted for this interest as an equity investment. On April 3, 2017, the Company acquired the remaining 60% ownership interest in CME. This acquisition did not materially impact the Company's consolidated financial results.

Transaction and other Acquisition-Related Costs

The Company incurred integration, restructuring and transaction costs relating to acquisitions in 2017, 2016 and 2015:

<u>(Millions of dollars)</u>	2017	2016	2015
Integration costs	\$ 237	\$ 192	\$ 95
Restructuring costs	85	526	271
Transaction costs	39	10	59

The Company's integration costs incurred in 2017, 2016 and 2015 were substantially associated with the CareFusion acquisition and a small portion of costs incurred in 2017 related to the pending Bard acquisition. The restructuring costs incurred in 2017, 2016 and 2015, which related to CareFusion and other portfolio rationalization initiatives, are further discussed in Note 11. Transaction costs, which primarily consisted of legal, advisory and other costs, were largely incurred in 2017 in connection with the pending agreement to acquire Bard and other portfolio rationalization initiatives. Transaction costs were also incurred in 2015 in connection with the CareFusion acquisition. The Company's integration, restructuring and transaction costs were primarily recorded as *Acquisitions and other restructurings*.

Discussion regarding the financing costs relating to the pending Bard acquisition and the CareFusion acquisition, which were recorded as *Interest expense*, are provided in Note 15.

Note 10 — Divestiture

Respiratory Solutions

On October 3, 2016, the Company sold a 50.1% controlling financial interest in its Respiratory Solutions business, a component of the Medical segment, to form a venture, Vyair Medical. The Company retained a 49.9% non-controlling interest in the new standalone entity. The buyer controls the operations and governance of the new entity. The Company accounts for its remaining interest in the new entity as an equity method investment and, beginning on January 1, 2017, records its share of the new entity's earnings or losses on a one-quarter lag to *Other income (expense), net*. The Company has agreed to various contract manufacturing and certain logistical and transition services agreements with the new entity for a period of up to two years after the sale. Assets and liabilities held for sale on the consolidated balance sheet at September 30, 2016 included assets and liabilities subject to this agreement of approximately \$578 million and \$189 million, respectively. The historical financial results for the Respiratory Solutions business, which included approximately \$822 million and \$417 million of revenues for years ended September 30, 2016 and 2015, respectively, have not been classified as a discontinued operation.

Note 11 — Business Restructuring Charges

In connection with the CareFusion acquisition and portfolio rationalization initiatives, the Company incurred restructuring costs which were recorded as *Acquisitions and other restructurings*. Additional disclosures regarding these restructuring activities and the related costs are provided in Notes 7, 9 and 10. Restructuring liability activity in 2017, 2016 and 2015 was as follows:

(Millions of dollars)	Employee Termination (A)	Other (B)	Total
Balance at September 30, 2014	\$ —	\$ —	\$ —
Assumed liability	19	—	19
Charged to expense	126	146	271
Cash payments	(74)	(20)	(94)
Non-cash settlements	—	(44)	(44)
Other adjustments	(9)	(81)	(91)
Balance at September 30, 2015	<u>\$ 62</u>	<u>\$ —</u>	<u>\$ 62</u>
Charged to expense	81	445	526
Cash payments	(76)	(72)	(148)
Non-cash settlements	—	(39)	(39)
Other adjustments	—	(332)	(332)
Balance at September 30, 2016	<u>\$ 67</u>	<u>\$ 2</u>	<u>\$ 69</u>
Charged to expense	27	58	85
Cash payments	(45)	(12)	(57)
Non-cash settlements	—	(9)	(9)
Other adjustments	—	(33)	(33)
Balance at September 30, 2017	<u><u>\$ 49</u></u>	<u><u>\$ 6</u></u>	<u><u>\$ 55</u></u>

- (A) Expenses in fiscal year 2016 included \$40 million relating to the CareFusion acquisition as well as \$13 million for employee termination costs resulting from the Company's transition of certain elements of its information technology function to an outsourced model as further disclosed below. Expenses in fiscal year 2015 were primarily related to the CareFusion acquisition.
- (B) In the fourth quarter of fiscal year 2015, the Company initiated a plan to transition certain elements of its global information technology function from Company-owned assets to an outsourced, cloud-based model. Restructuring expenses recorded in 2015 included non-cash asset impairment charges relating to assets held for sale as a result of this plan. In the fourth quarter of fiscal year 2016, the Company expanded the scope of this plan to include the transition of certain business information systems assets to a third-party and as a result, the Company recorded a \$214 million non-cash charge to recognize the impairment of capitalized internal-use software assets held for sale. Expenses in 2016 also included non-cash impairment charges of \$81 million, after-tax, relating to the Company's disposition of certain non-core businesses, including the Company's sale of a majority interest in its Respiratory Solutions business during the first quarter of fiscal year 2017, which is further discussed in Note 10.

Note 12 — Intangible Assets

Intangible assets at September 30 consisted of:

(Millions of dollars)	2017		2016	
	Gross Carrying Amount	Accumulated Amortization	Gross Carrying Amount	Accumulated Amortization
<i>Amortized intangible assets</i>				
Customer relationships	\$ 3,393	\$ 564	\$ 3,360	\$ 339
Developed technology	3,508	1,029	3,409	754
Product rights	131	54	125	43
Trademarks	408	65	405	45
Patents and other	370	274	349	254
Amortized intangible assets	<u>\$ 7,811</u>	<u>\$ 1,986</u>	<u>\$ 7,648</u>	<u>\$ 1,435</u>
<i>Unamortized intangible assets</i>				
Acquired in-process research and development	\$ 67		\$ 66	
Trademarks	2		2	
Unamortized intangible assets	<u>\$ 69</u>		<u>\$ 68</u>	

Intangible amortization expense was \$533 million, \$552 million and \$346 million in 2017, 2016 and 2015, respectively. The estimated aggregate amortization expense for the fiscal years ending September 30, 2018 to 2022 are as follows: 2018 — \$526 million; 2019 — \$519 million; 2020 — \$518 million; 2021 — \$514 million; 2022 — \$503 million.

The following is a reconciliation of goodwill by business segment:

(Millions of dollars)	Medical	Life Sciences	Total
Goodwill as of September 30, 2015	\$ 6,807	\$ 730	\$ 7,537
Divestiture (A)	(32)	—	(32)
Purchase accounting adjustments (B)	(79)	1	(78)
Currency translation	(8)	—	(8)
Goodwill as of September 30, 2016	\$ 6,688	\$ 731	\$ 7,419
Acquisitions (C)	119	24	143
Divestiture (A)	(25)	—	(25)
Purchase accounting adjustments	4	—	4
Currency translation	16	6	22
Goodwill as of September 30, 2017	<u>\$ 6,802</u>	<u>\$ 761</u>	<u>\$ 7,563</u>

- (A) Represents goodwill derecognized upon the Company's sale of a 50.1% controlling financial interest in the Respiratory Solutions business, as further discussed in Note 10.
- (B) Primarily represents \$94 million resulting from adjustments to the deferred tax liability accounts recorded upon the Company's acquisition of CareFusion.
- (C) Represents goodwill recognized upon the Company's acquisitions during the current year. Such acquisitions were not material individually or in the aggregate.

Note 13 — Derivative Instruments and Hedging Activities

The Company uses derivative instruments to mitigate certain exposures. The Company does not enter into derivative financial instruments for trading or speculative purposes. The effects these derivative instruments and hedged items have on financial position, financial performance, and cash flows are provided below.

Foreign Currency Risks and Related Strategies

The Company has foreign currency exposures throughout Europe, Greater Asia, Canada and Latin America. Transactional currency exposures that arise from entering into transactions, generally on an intercompany basis, in non-hyperinflationary countries that are denominated in currencies other than the functional currency are mitigated primarily through the use of forward contracts and currency options. Hedges of the transactional foreign exchange exposures resulting primarily from intercompany payables and receivables are undesignated hedges. As such, the gains or losses on these instruments are recognized immediately in income. These gains and losses are largely offset by gains and losses on the underlying hedged items, as well as the hedging costs associated with the derivative instruments. The net amounts in 2017, 2016 and 2015, which are recognized in *Other income (expense), net*, were immaterial to the Company's consolidated financial results.

The total notional amounts of the Company's outstanding foreign exchange contracts as of September 30, 2017 and 2016 were \$2.5 billion and \$2.3 billion, respectively.

In order to mitigate foreign currency exposure relating to its investments in certain foreign subsidiaries, the Company has designated \$2 billion of euro-denominated debt, issued during the first and third quarters of fiscal year 2017, as net investment hedges. Accordingly, net gains or losses relating to this debt, which are attributable to changes in the euro to U.S. dollar spot exchange rate, are recorded as accumulated foreign currency translation in *Other comprehensive income (loss)*. Recognition of hedge ineffectiveness into earnings will occur if the notional amount of the euro-denominated debt no longer matches the portion of the net investments in foreign subsidiaries which underlie the hedges. The Company's balance of *Accumulated other comprehensive income (loss)* as of September 30, 2017 included net losses relating to these net investment

hedges of \$159 million. Additional disclosures regarding the Company's issuances of the euro-denominated debt in fiscal year 2017 are provided in Note 15.

Interest Rate Risks and Related Strategies

The Company's primary interest rate exposure results from changes in U.S. dollar interest rates. The Company's policy is to manage interest cost using a mix of fixed and variable rate debt. The Company periodically uses interest rate swaps to manage such exposures. Under these interest rate swaps, the Company exchanges, at specified intervals, the difference between fixed and floating interest amounts calculated by reference to an agreed-upon notional principal amount. These swaps are designated as either fair value or cash flow hedges.

For interest rate swaps designated as fair value hedges (i.e., hedges against the exposure to changes in the fair value of an asset or a liability or an identified portion thereof that is attributable to a particular risk), changes in the fair value of the interest rate swaps offset changes in the fair value of the fixed rate debt due to changes in market interest rates.

Changes in the fair value of the interest rate swaps designated as cash flow hedges (i.e., hedging the exposure to variability in expected future cash flows that is attributable to a particular risk) are offset by amounts recorded in *Other comprehensive income (loss)*. If interest rate derivatives designated as cash flow hedges are terminated, the balance in *Accumulated other comprehensive income (loss)* attributable to those derivatives is reclassified into earnings over the remaining life of the hedged debt. The net realized loss related to terminated interest rate swaps expected to be reclassified and recorded in *Interest expense* within the next 12 months is \$4 million, net of tax.

At September 30, 2016, the total notional value of the Company's outstanding forward starting interest rate swaps, which were entered into to mitigate the Company's exposure to interest rate risk and were designated as cash flow hedges, was \$500 million. In the third quarter of 2017, prior to its issuance of senior unsecured U.S. notes during same quarter, the Company entered into additional forward starting interest rate swaps, which were also designated as cash flow hedges, with a notional amount of \$1.75 billion. The Company's recognition of unrealized and realized amounts, including net realized losses recognized upon the termination of all of the Company's outstanding interest rate hedges in the third quarter of 2017 concurrently with its issuance of the senior unsecured notes noted above, resulted in the Company's recognition of a net gain in other comprehensive income relating to interest rate hedges during the year ended September 30, 2017. Additional disclosures regarding the Company's issuance of notes are provided in Note 15 and additional disclosures regarding the net gain recognized on terminated cash flow hedges during 2017 are provided in Note 3.

The total notional amount of the Company's outstanding interest rate swaps designated as fair value hedges was \$375 million at September 30, 2017 and 2016. The outstanding swaps represent fixed-to-floating interest rate swap agreements the Company entered into to convert the interest payments on \$375 million of the Company's 3.125% notes due 2021 from the fixed rate to a floating interest rate based on LIBOR. Changes in the fair value of the interest rate swaps offset changes in the fair value of the fixed rate debt. The (losses) gains recorded on these fair value hedges, which were offset by (gains) losses recorded to the underlying debt instruments, were immaterial to the Company's consolidated financial results.

Other Risk Exposures

The Company purchases resins, which are oil-based components used in the manufacture of certain products. Significant increases in world oil prices that lead to increases in resin purchase costs could impact future operating results. From time to time, the Company has managed price risks associated with these commodity purchases through commodity derivative forward contracts. The Company had no outstanding commodity derivative forward contracts at September 30, 2017 and 2016.

Effects on Consolidated Balance Sheets

The location and amounts of derivative instrument fair values in the consolidated balance sheet are segregated below between designated, qualifying hedging instruments and ones that are not designated for hedge accounting.

(Millions of dollars)	September 30, 2017	September 30, 2016
Asset derivatives-designated for hedge accounting		
Interest rate swaps	\$ 7	\$ 23
Asset derivatives-undesignated for hedge accounting		
Forward exchange contracts	8	3
Total asset derivatives (A)	<u>\$ 15</u>	<u>\$ 25</u>
Liability derivatives-designated for hedge accounting		
Interest rate swaps	—	18
Liability derivatives-undesignated for hedge accounting		
Forward exchange contracts	7	13
Total liability derivatives (B)	<u>\$ 7</u>	<u>\$ 31</u>

(A) All asset derivatives are included in *Prepaid expenses and other*.

(B) All liability derivatives are included in *Accrued expenses*.

Effects on Consolidated Statements of Income

The amounts recognized in other comprehensive income during 2017, 2016 and 2015 relating to cash flow hedges were not material to the Company's consolidated financial results. The Company's designated derivative instruments are highly effective. As such, there were no gains or losses, related to hedge ineffectiveness or amounts excluded from hedge effectiveness testing, recognized immediately in income relative to derivative contracts outstanding in the periods presented.

Note 14 — Financial Instruments and Fair Value Measurements

Recurring Fair Value Measurements

The fair values of financial instruments, including those not recognized on the statement of financial position at fair value, carried at September 30, 2017 and 2016 are classified in accordance with the fair value hierarchy in the following table:

(Millions of dollars)	Basis of Fair Value Measurement							
	Total		Quoted Prices in Active Markets for Identical Assets (Level 1)		Significant Other Observable Inputs (Level 2)		Significant Unobservable Inputs (Level 3)	
	2017	2016	2017	2016	2017	2016	2017	2016
<u>Assets</u>								
Institutional money market investments	\$ 2,026	\$ 224	\$ 2,026	\$ 224	\$ —	\$ —	\$ —	\$ —
Interest rate swaps	7	23	—	—	7	23	—	—
Forward exchange contracts	8	3	—	—	8	3	—	—
Total Assets	<u>\$ 2,042</u>	<u>\$ 249</u>	<u>\$ 2,026</u>	<u>\$ 224</u>	<u>\$ 15</u>	<u>\$ 25</u>	<u>\$ —</u>	<u>\$ —</u>
<u>Liabilities</u>								
Interest rate swaps	\$ —	\$ 18	\$ —	\$ —	\$ —	\$ 18	\$ —	\$ —
Forward exchange contracts	7	13	—	—	7	13	—	—
Contingent consideration liabilities	13	54	—	—	—	—	13	54
Total Liabilities	<u>\$ 20</u>	<u>\$ 86</u>	<u>\$ —</u>	<u>\$ —</u>	<u>\$ 7</u>	<u>\$ 31</u>	<u>\$ 13</u>	<u>\$ 54</u>

The Company's institutional money market accounts permit daily redemption and the fair values of these investments are based upon the quoted prices in active markets provided by the holding financial institutions. The Company's remaining cash equivalents were \$12.153 billion and \$1.317 billion at September 30, 2017 and 2016, respectively. Short-term investments are held to their maturities and are carried at cost, which approximates fair value. The cash equivalents consist of liquid investments with a maturity of three months or less and the short-term investments consist of instruments with maturities greater than three months and less than one year.

The Company measures the fair value of forward exchange contracts and interest rate swaps based upon the present value of expected future cash flows using market-based observable inputs including credit risk, interest rate yield curves, foreign currency spot prices and forward prices.

Long-term debt is recorded at amortized cost. The fair value of long-term debt is measured based upon quoted prices in active markets for similar instruments, which are considered Level 2 inputs in the fair value hierarchy. The fair value of long-term debt was \$19.2 billion and \$11.3 billion at September 30, 2017 and 2016, respectively. The fair value of the current portion of long-term debt was \$206 million and \$798 million at September 30, 2017 and 2016, respectively.

The contingent consideration liabilities were recognized as part of the consideration transferred by the Company for certain acquisitions. The fair values of the contingent consideration liabilities were estimated using probability-weighted discounted cash flow models that were based upon the probabilities assigned with regard to achievement of the contingent events. The estimated fair values of the contingent consideration liabilities are remeasured each reporting period based upon increases or decreases in the probability of the contingent payments. The decrease to the total contingent consideration liability in fiscal year 2017 is primarily

attributable to a \$40 million payment of a contingent consideration liability recorded in connection with a previously closed acquisition.

The Company's policy is to recognize any transfers into fair value measurement hierarchy levels and transfers out of levels at the beginning of each reporting period. There were no transfers in and out of Level 1, Level 2 or Level 3 measurements for the years ending September 30, 2017 and 2016.

Nonrecurring Fair Value Measurements

In fiscal year 2016, the Company recorded a charge of \$214 million to impair capitalized internal-use software assets held for sale as a result of the Company's transition of certain elements of its information technology infrastructure to an outsourced model. Impairment charges recorded in 2015 of \$72 million primarily related to information technology assets held for sale as a result of these same transition efforts. Also in fiscal year 2016, the Company recorded losses of \$81 million on the held for sale assets of certain non-core businesses. The amounts recognized in 2016 and 2015 were recorded to *Acquisitions and other restructurings* to adjust the carrying amount of assets held for sale to an estimate of the assets' fair values, less the estimated costs to sell these assets. The fair values of the assets adjusted in 2016 and 2015 were estimated, based upon a market participant's perspective, using a market approach and Level 2 inputs, including quoted prices for similar assets.

Concentration of Credit Risk

The Company maintains cash deposits in excess of government-provided insurance limits. Such cash deposits are exposed to loss in the event of nonperformance by financial institutions. Substantially all of the Company's trade receivables are due from public and private entities involved in the healthcare industry. Due to the large size and diversity of the Company's customer base, concentrations of credit risk with respect to trade receivables are limited. The Company does not normally require collateral. The Company is exposed to credit loss in the event of nonperformance by financial institutions with which it conducts business. However, this loss is limited to the amounts, if any, by which the obligations of the counterparty to the financial instrument contract exceed the obligations of the Company. The Company also minimizes exposure to credit risk by dealing with a diversified group of major financial institutions.

The Company continually evaluates its accounts receivables for potential collection risks particularly those resulting from sales to government-owned or government-supported healthcare facilities in certain countries as payment may be dependent upon the financial stability and creditworthiness of those countries' national economies. The Company continually evaluates all governmental receivables for potential collection risks associated with the availability of government funding and reimbursement practices. The Company believes the current reserves related to all governmental receivables are adequate and that this concentration of credit risk will not have a material adverse impact on its financial position or liquidity.

Note 15 — Debt**Short-term debt**

Short-term debt at September 30 consisted of:

(Millions of dollars)	2017	2016
Commercial paper borrowings	\$ —	\$ 200
Current portion of long-term debt		
4.900% Notes due April 15, 2018	200	—
1.450% Notes due May 15, 2017 (A)	—	300
1.750% Notes due November 8, 2016	—	500
Other	3	1
Total short-term debt	<u>\$ 203</u>	<u>\$ 1,001</u>

(A) All or a portion of the aggregate principal amount outstanding was repurchased during the first quarter of 2017, as further discussed below.

The weighted average interest rates for short-term debt were 4.90% and 1.49% at September 30, 2017 and 2016, respectively.

Long-term debt

Long-Term Debt at September 30 consisted of:

(Millions of dollars)		2017	2016
1.800% Notes due December 15, 2017	(A) (C)	—	1,248
4.900% Notes due April 15, 2018		—	201
5.000% Notes due May 15, 2019	(A) (C)	—	498
2.133% Notes due June 6, 2019	(B)	723	—
0.368% Notes due June 6, 2019	(B)	823	—
6.375% Notes due August 1, 2019	(A) (C)	—	776
2.675% Notes due December 15, 2019	(A)	1,121	1,245
2.404% Notes due June 5, 2020	(B)	996	—
3.250% Notes due November 12, 2020		698	698
3.125% Notes due November 8, 2021		1,003	1,018
2.894% Notes due June 6, 2022	(B)	1,791	—
Floating Rate Notes due June 6, 2022	(B)	497	—
1.000% Notes due December 15, 2022	(D)	586	—
3.300% Notes due March 1, 2023	(C)	296	304
3.875% Notes due May 15, 2024	(A) (C)	182	417
3.363% Notes due June 6, 2024	(B)	1,736	—
3.734% Notes due December 15, 2024	(A)	1,367	1,740
1.900% Notes due December 15, 2026	(D)	585	—
3.700% Notes due June 6, 2027	(B)	2,381	—
7.000% Debentures due August 1, 2027		166	168
6.700% Debentures due August 1, 2028		164	167
6.000% Notes due May 15, 2039		246	246
5.000% Notes due November 12, 2040		296	297
4.875% Notes due May 15, 2044	(C)	331	333
4.685% Notes due December 15, 2044		1,189	1,190
4.669% Notes due June 6, 2047	(B)	1,484	—
Other long-term debt		3	4
Total Long-Term Debt		<u>\$ 18,667</u>	<u>\$ 10,550</u>

- (A) All or a portion of the aggregate principal amount outstanding was repurchased during the first quarter of 2017, as further discussed below.
- (B) Notes issued during the third quarter of fiscal year 2017, as further discussed below.
- (C) All or a portion of the aggregate principal amount outstanding was redeemed during the third quarter of 2017, as further discussed below.
- (D) Notes issued during the first quarter of fiscal year 2017, as further discussed below.

The aggregate annual maturities of long-term debt including interest during the fiscal years ending September 30, 2018 to 2022 are as follows: 2018 — \$823 million; 2019 — \$2.2 billion; 2020 — \$2.8 billion; 2021 — \$1.2 billion; 2022 — \$3.8 billion.

Other current credit facilities

In January 2016, the Company replaced an existing \$1 billion syndicated credit facility with a \$1.5 billion syndicated credit facility. During the first quarter of fiscal year 2017, the Company extended the expiration date of this credit facility to January 2022 from the original expiration date of January 2021. There were no

borrowings outstanding under this credit facility at September 30, 2017. The credit facility, under which the Company may issue up to \$100 million in letters of credit, can be used for general corporate purposes. It includes a provision that enables BD, subject to additional commitments made by the lenders, to access up to an additional \$500 million in financing through the facility for a maximum aggregate commitment of \$2 billion. The credit facility includes a single financial covenant that requires BD to maintain an interest expense coverage ratio of not less than 4-to-1 for the most recent four consecutive fiscal quarters. The Company was in compliance with this covenant as of September 30, 2017. The Company will replace this existing syndicated credit facility with a five-year senior unsecured revolving credit facility upon the closing of the Bard acquisition, as further discussed below. In addition, the Company has informal lines of credit outside of the United States.

First Quarter Fiscal Year 2017 Debt-Related Transactions

In December 2016, the Company issued euro-denominated debt consisting of 500 million euros of 1.000% notes and 500 million euros of 1.900% notes. The Company used the net proceeds from this long-term debt offering, together with other sources of liquidity, to fund the Company's repurchase of certain of its senior notes outstanding. Under this cash tender offer, the Company repurchased all or a portion of the aggregate principal amounts of certain of its long-term notes outstanding, totaling \$1.689 billion, at an aggregate market price of \$1.764 billion. The carrying value of these long-term notes was \$1.727 billion, and the Company recognized a loss on this debt extinguishment of \$42 million, which was recorded in December 2016 as *Other income (expense), net*, on the Company's consolidated statements of income.

Third Quarter Fiscal Year 2017 Debt-Related Transactions

In connection with the Company's agreement to acquire Bard, as previously discussed in Note 9, the Company's capital structure was impacted in the third quarter by the debt-related transactions discussed below.

- The Company entered into a three-year senior unsecured term loan facility of \$2.25 billion. The proceeds from this facility may only be used to fund a portion of the cash consideration for the Bard acquisition, as well as the fees and expenses incurred in connection with this acquisition, which is expected to close in the fourth calendar quarter of 2017.
- The Company also entered into a five-year senior unsecured revolving credit facility that will provide borrowing of up to \$2.25 billion when the facility becomes effective upon the closing of the Bard acquisition. The facility, which will expire in May 2022, will replace the syndicated credit facility the Company currently has in place, as discussed above. The Company intends to use the new revolving facility to fund general corporate needs and to redeem, repurchase or defease certain of Bard's outstanding senior unsecured notes that will be assumed upon the closing of the acquisition.
- The Company issued senior unsecured U.S. notes with an aggregate principal amount of \$9.675 billion. If the Company's acquisition of Bard does not close by April 23, 2018, or if the agreement to acquire Bard is terminated prior to this date, the Company will be required to redeem all of the senior unsecured U.S. notes issued as detailed above, except for the notes which are due in 2019. The notes would be redeemed at a special mandatory redemption price equal to 101% of their aggregate principal amount, plus accrued and unpaid interest to, but excluding, the redemption date.
- The Company issued Euro-denominated debt consisting of 700 million euros of 0.368% Notes due June 6, 2019.
- The Company redeemed all or a portion of the aggregate principal amounts of certain of its long-term senior notes outstanding, totaling \$1.717 billion, at an aggregate market price of \$1.776 billion. The carrying value of these long-term notes was \$1.745 billion and the Company recognized a loss on this debt extinguishment of \$31 million, which was recorded in June 2017 as *Other income (expense), net*, on the Company's consolidated statements of income.

- Upon securing the permanent financing arrangements discussed above, an agreement for \$15.7 billion of fully committed bridge financing that was entered into concurrently with the execution of the agreement to acquire Bard was terminated.

Also in connection with the Company's agreement to acquire Bard, the Company commenced offers to exchange certain outstanding notes issued by Bard for a like-amount of new notes to be issued by the Company. The offers are conditioned upon the closing of the Bard acquisition and the expiration of these offers will be extended until the acquisition closes. The aggregate principal amounts of Bard notes which have been validly tendered for notes issued by the Company since the offers were commenced in May are provided below.

(Millions of dollars)	Aggregate Principal Amount	Principal Amount Accepted for Exchange
4.400% Notes due January 15, 2021	\$ 500	\$ 428
3.000% Notes due May 15, 2026	500	470
6.700% Notes due December 1, 2026	150	137
Total	<u>\$ 1,150</u>	<u>\$ 1,035</u>

Capitalized interest

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

(Millions of dollars)	2017	2016	2015
Charged to operations	\$ 521	\$ 388	\$ 371
Capitalized	32	30	30
Total interest costs	<u>\$ 553</u>	<u>\$ 418</u>	<u>\$ 401</u>
Interest paid, net of amounts capitalized	<u>\$ 435</u>	<u>\$ 392</u>	<u>\$ 313</u>

Note 16 — Income Taxes

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

(Millions of dollars)	2017	2016	2015
Current:			
Federal	\$ (230)	\$ 312	\$ 50
State and local, including Puerto Rico	(20)	17	15
Foreign	200	286	252
	<u>\$ (50)</u>	<u>\$ 616</u>	<u>\$ 318</u>
Deferred:			
Domestic	\$ (64)	\$ (441)	\$ (238)
Foreign	(10)	(78)	(37)
	<u>(74)</u>	<u>(519)</u>	<u>(274)</u>
Income tax provision	<u>\$ (124)</u>	<u>\$ 97</u>	<u>\$ 44</u>

The components of *Income Before Income Taxes* for the years ended September 30 consisted of:

(Millions of dollars)	2017	2016	2015
Domestic, including Puerto Rico	\$ (386)	\$ (232)	\$ (408)
Foreign	1,362	1,306	1,147
Income Before Income Taxes	<u>\$ 976</u>	<u>\$ 1,074</u>	<u>\$ 739</u>

The table below summarizes the gross amounts of unrecognized tax benefits without regard to reduction in tax liabilities or additions to deferred tax assets and liabilities if such unrecognized tax benefits were settled. The Company believes it is reasonably possible that certain audits will close within the next twelve months but no significant increases or decreases in the amount of the unrecognized tax benefits are expected to result.

(Millions of dollars)	2017	2016	2015
Balance at October 1	\$ 469	\$ 593	\$ 206
Increase due to acquisitions	—	—	326
Increase due to current year tax positions	41	81	62
Increase due to prior year tax positions	19	10	14
Decreases due to prior year tax positions	(30)	(3)	(2)
Decrease due to settlements with tax authorities	(145)	(147)	(10)
Decrease due to lapse of statute of limitations	(5)	(65)	(3)
Balance at September 30	<u>\$ 349</u>	<u>\$ 469</u>	<u>\$ 593</u>

Upon the Company's acquisition of CareFusion, the Company became a party to a tax matters agreement with Cardinal Health resulting from Cardinal Health's spin-off of CareFusion in fiscal year 2010. Under the tax matters agreement, the Company is obligated to indemnify Cardinal Health for certain tax exposures and transaction taxes prior to CareFusion's spin-off from Cardinal Health. The indemnification payable is approximately \$134 million at September 30, 2017 and is included in *Deferred Income Taxes and Other* on the consolidated balance sheet.

At September 30, 2017, 2016 and 2015, there are \$415 million, \$478 million and \$620 million of unrecognized tax benefits that if recognized, would affect the effective tax rate. During the fiscal years ended September 30, 2017, 2016 and 2015, the Company reported interest and penalties associated with unrecognized tax benefits of \$57 million, \$(38) million and \$5 million on the consolidated statements of income as a component of *Income tax provision*.

The Company conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of tax jurisdictions. While the IRS has completed its audit for the BD legacy fiscal year 2014 and combined company fiscal year 2016, the IRS has recently commenced the audit for the CareFusion legacy fiscal year 2014 and short period 2015. For the BD legacy business, all years are effectively settled with the exception of 2015 for which the Company believes it is adequately reserved for any potential exposures. With the exception of the CareFusion legacy fiscal year 2010 audit, all other periods are at various stages of appeals or protests. For the Company's other major tax jurisdictions where it conducts business, the Company's tax years are generally open after 2011.

Deferred income taxes at September 30 consisted of:

(Millions of dollars)	2017		2016	
	Assets	Liabilities	Assets	Liabilities
Compensation and benefits	\$ 618	\$ —	\$ 720	\$ —
Property and equipment	—	244	—	164
Intangibles	—	1,584	—	1,571
Loss and credit carryforwards	1,098	—	1,101	—
Other	531	164	783	664
	<u>2,247</u>	<u>1,992</u>	<u>2,604</u>	<u>2,399</u>
Valuation allowance	(1,032)	—	(997)	—
Net (A)	<u>\$ 1,216</u>	<u>\$ 1,992</u>	<u>\$ 1,606</u>	<u>\$ 2,399</u>

(A) Net deferred tax assets are included in *Other Assets* and net deferred tax liabilities are included in *Deferred Income Taxes and Other*.

Deferred tax assets and liabilities are netted on the balance sheet by separate tax jurisdictions. Deferred taxes are not provided on undistributed earnings of foreign subsidiaries that are indefinitely reinvested. At September 30, 2017, the cumulative amount of such undistributed earnings indefinitely reinvested outside the United States was \$9.6 billion. Determining the tax liability that would arise if these earnings were remitted is not practicable. Deferred taxes are provided for earnings outside the United States when those earnings are not considered indefinitely reinvested.

Generally, deferred tax assets have been established as a result of net operating losses and credit carryforwards with expiration dates from 2018 to an unlimited expiration date. Valuation allowances have been established as a result of an evaluation of the uncertainty associated with the realization of certain deferred tax assets on these losses and credit carryforwards. The valuation allowance for 2017 is primarily the result of foreign losses due to the Company's global re-organization of its foreign entities and these generally have no expiration date. Valuation allowances are also maintained with respect to deferred tax assets for certain federal and state carryforwards that may not be realized and that principally expire in 2021.

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

	2017	2016	2015
Federal statutory tax rate	35.0 %	35.0%	35.0%
State and local income taxes, net of federal tax benefit	(2.6)	1.5	(3.6)
Effect of foreign and Puerto Rico earnings and foreign tax credits	(40.8)	(23.7)	(24.5)
Effect of Research Credits and Domestic Production Activities	(2.7)	(4.4)	(1.6)
Effect of change in accounting for excess tax benefit relating to share-based compensation (see Note 2)	(7.9)	—	—
Other, net	6.3	0.7	0.6
Effective income tax rate	<u>(12.7)%</u>	<u>9.1%</u>	<u>5.9%</u>

The approximate amounts of tax reductions related to tax holidays in various countries in which the Company does business were \$144 million, \$121 million and \$103 million, in 2017, 2016 and 2015, respectively. The benefit of the tax holiday on diluted earnings per share was approximately \$0.64, \$0.56, and \$0.48 for fiscal years 2017, 2016 and 2015, respectively. The tax holidays expire at various dates through 2026.

The Company made income tax payments, net of refunds, of \$265 million in 2017, \$218 million in 2016 and \$240 million in 2015.

Note 17 — Sales-Type Leases and Financing Receivables

In April 2017, in conjunction with the implementation of a new “go-to-market” business model for the Company’s U.S. dispensing business within the Medication Management Solutions (“MMS”) unit of the Medical segment, the Company amended the terms of certain customer leases for dispensing equipment within the MMS unit. The modification provided customers the ability to reduce its dispensing asset base via a return provision, resulting in a more flexible lease term. Prior to the modification, these leases were accounted for as sales-type leases in accordance with Accounting Standards Codification Topic 840, “Leases”, as the non-cancellable lease term of 5 years exceeded 75% of the equipment’s estimated useful life and the present value of the minimum lease payments exceeded 90% of the equipment’s fair value. As a result of the lease modification, the Company was required to reassess the classification of the leases due to the amended lease term. Accordingly, most amended lease contracts were classified as operating leases beginning in April 2017. The change in lease classification resulted in a pre-tax charge to earnings in fiscal year 2017 of \$748 million, which was recorded in *Other operating (income) expense, net*, relating to the derecognition of the net investment in sales-type leases of \$1.065 billion, partially offset by the recognition of the underlying leased assets, as *Property, Plant and Equipment, Net* on the Company’s balance sheet, as of the effective date of \$317 million. Beginning April 1, 2017, revenue associated with these modified contracts is recognized on a straight-line basis over the remaining lease term, along with depreciation on the reinstated leased assets.

The Company’s consolidated financial results in 2017 were not materially impacted by the financing receivables remaining subsequent to the lease modification discussed above.

Note 18 — Supplemental Financial Information

Other Income (Expense), Net

<u>(Millions of dollars)</u>	<u>2017</u>	<u>2016</u>	<u>2015</u>
Losses on debt extinguishment	\$ (73)	\$ —	\$ —
Share of Vyaire Medical venture results, net of income from transition services agreements	(3)	—	—
Other equity investment income	3	8	9
Losses on undesignated foreign exchange derivatives, net	(11)	(3)	—
Gains on previously held investments	24	—	9
Other	3	7	3
Other (expense) income, net	<u>\$ (57)</u>	<u>\$ 11</u>	<u>\$ 21</u>

Additional disclosures regarding the losses recognized on the extinguishment of debt are provided in Note 15. Additional disclosures regarding the Company’s divestiture of the Respiratory Solutions business and the Vyaire Medical venture formed with this business are provided in Note 10. The Company recognized an acquisition-date accounting gain related to a previously-held equity method investment in an entity that the Company acquired during the third quarter of fiscal year 2017. The Company also recognized such a gain in 2015, when it acquired another entity in which an equity method investment was held prior to the acquisition date.

Trade Receivables, Net

The amounts recognized in 2017, 2016 and 2015 relating to allowances for doubtful accounts and cash discounts, which are netted against trade receivables, are provided in the following table:

(Millions of dollars)	Allowance for Doubtful Accounts	Allowance for Cash Discounts	Total
Balance at September 30, 2014	\$ 30	\$ 12	\$ 42
Additions charged to costs and expenses	33	47	80
Deductions and other	(11) (A)	(50)	(61)
Balance at September 30, 2015	\$ 53	\$ 9	\$ 62
Additions charged to costs and expenses	23	37	60
Deductions and other	(14) (A)	(40)	(55)
Balance at September 30, 2016	\$ 61	\$ 6	\$ 67
Additions charged to costs and expenses	25	43	68
Deductions and other	(32) (A)	(45)	(76)
Balance at September 30, 2017	\$ 54	\$ 4	\$ 58

(A) Accounts written off.

Inventories

Inventories at September 30 consisted of:

(Millions of dollars)	2017	2016
Materials	\$ 313	\$ 316
Work in process	271	274
Finished products	1,234	1,129
	\$ 1,818	\$ 1,719

Property, Plant and Equipment, Net

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

(Millions of dollars)	2017	2016
Land	\$ 146	\$ 151
Buildings	2,496	2,397
Machinery, equipment and fixtures	6,584	5,749
Leasehold improvements	163	121
	9,389	8,419
Less accumulated depreciation and amortization	4,752	4,518
	\$ 4,638	\$ 3,901

The increase in Machinery, equipment and fixtures included \$317 million of assets recognized as *Property, Plant and Equipment, Net* on the Company's balance sheet, resulting from a modification to the Company's dispensing equipment lease contracts with customers. Additional disclosures regarding this modification are provided in Note 17.

SUPPLEMENTARY QUARTERLY DATA (UNAUDITED)

Millions of dollars, except per share amounts

	2017				
	1 st	2 nd	3 rd	4 th	Year
Revenues	\$ 2,922	\$ 2,969	\$ 3,035	\$ 3,166	\$ 12,093
Gross Profit	1,452	1,432	1,504	1,554	5,942
Net Income (Loss)	562	344	(132)	327	1,100
Earnings per Share:					
Basic	2.64	1.61	(0.75)	1.27	4.70
Diluted	2.58	1.58	(0.75)	1.24	4.60
	2016				
	1 st	2 nd	3 rd	4 th	Year
Revenues	\$ 2,986	\$ 3,067	\$ 3,198	\$ 3,231	\$ 12,483
Gross Profit	1,408	1,484	1,547	1,552	5,991
Net Income	229	338	390	19	976
Earnings per Share:					
Basic	1.08	1.59	1.83	0.09	4.59
Diluted	1.06	1.56	1.80	0.09	4.49

Certain quarterly amounts may not add to the year-to-date totals due to rounding. Earnings per share amounts are calculated from the underlying whole-dollar amounts. As of May 2017, the weighted average common shares used in the computations of basic and diluted earnings per share reflect shares issued in anticipation of the pending acquisition of Bard. Additional disclosures regarding this issuance of shares and the pending acquisition are provided in Notes 3 and 9.

Item 9. Changes in and Disagreements With Accountants on Accounting and Financial Disclosure.

None.

Item 9A. Controls and Procedures.

An evaluation was conducted by BD's management, with the participation of BD's Chief Executive Officer and Chief Financial Officer, of the effectiveness of the design and operation of BD's disclosure controls and procedures (as defined in Rule 13a-15(e) under the Securities Exchange Act of 1934) as of September 30, 2017. Based upon that evaluation, the Chief Executive Officer and Chief Financial Officer concluded that the design and operation of these disclosure controls and procedures were, as of the end of the period covered by this report, effective and designed to ensure that material information relating to BD and its consolidated subsidiaries would be made known to them by others within these entities. There were no changes in our internal control over financial reporting during the fiscal quarter ended September 30, 2017 identified in connection with the above-referenced evaluation that have materially affected, or are reasonably likely to materially affect, BD's internal control over financial reporting.

Management's Report on Internal Control Over Financial Reporting and the Report of Independent Registered Public Accounting Firm are contained in Item 8, Financial Statements and Supplementary Data, and are incorporated herein by reference.

Item 9B. Other Information.

None.

PART III

Item 10. *Directors, Executive Officers and Corporate Governance.*

The information relating to directors and the Audit Committee of the BD Board of Directors required by this item will be contained under the captions “Proposal 1. Election of Directors” and “Board of Directors - Committee Membership and Function - Audit Committee” in a definitive proxy statement involving the election of directors, which the registrant will file with the SEC not later than 120 days after September 30, 2017 (the “2018 Proxy Statement”), and such information is incorporated herein by reference.

The information relating to executive officers required by this item is included herein in Part I under the caption “Executive Officers of the Registrant.”

Certain other information required by this item will be contained under the captions “Ownership of BD Common Stock - Section 16(a) beneficial ownership reporting compliance”, “Corporate Governance - Director nomination process” and Corporate Governance - Code of Conduct” in BD’s 2018 Proxy Statement, and such information is incorporated herein by reference.

Item 11. *Executive Compensation.*

The information required by this item will be contained under the captions “Board of Directors - Non-Management Directors’ Compensation,” “Compensation Discussion and Analysis,” “Report of the Compensation and Management Development Committee,” “Compensation of Named Executive Officers” and “Board of Directors - Committee membership and function - Compensation and Management Development Committee” in BD’s 2018 Proxy Statement, and such information is incorporated herein by reference.

Item 12. *Security Ownership of Certain Beneficial Owners and Management and Related Stockholder Matters.*

The information required by this item will be contained under the caption “Ownership of BD Common Stock” in BD’s 2018 Proxy Statement, and such information is incorporated herein by reference.

Item 13. *Certain Relationships and Related Transactions, and Director Independence.*

The information required by this item will be contained under the caption “Corporate Governance - Director Independence; Policy Regarding Related Person Transactions” in BD’s 2018 Proxy Statement, and such information is incorporated herein by reference.

Item 14. *Principal Accounting Fees and Services.*

The information required by this item will be contained under the caption “Proposal 2. Ratification of Selection of Independent Registered Public Accounting Firm” in BD’s 2018 Proxy Statement, and such information is incorporated herein by reference.

PART IV

Item 15. Exhibits, Financial Statement Schedules.

(a)(1) *Financial Statements*

The following consolidated financial statements of BD are included in Item 8 of this report:

- Reports of Independent Registered Public Accounting Firm
- Consolidated Statements of Income — Years ended September 30, 2017, 2016 and 2015
- Consolidated Statements of Comprehensive Income — Years ended September 30, 2017, 2016 and 2015
- Consolidated Balance Sheets — September 30, 2017 and 2016
- Consolidated Statements of Cash Flows — Years ended September 30, 2017, 2016 and 2015
- Notes to Consolidated Financial Statements

(2) *Financial Statement Schedules*

See Note 18 to the Consolidated Financial Statements included in Item 8, Financial Statements and Supplementary Data.

(3) *Exhibits*

See the Exhibit Index beginning on page 99 hereof for a list of all management contracts, compensatory plans and arrangements required by this item, and all other Exhibits filed or incorporated by reference as a part of this report.

SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) 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.

BECTON, DICKINSON AND COMPANY

By: /s/ GARY DEFazio
Gary DeFazio
Senior Vice President and Corporate Secretary

Dated: November 22, 2017

Pursuant to the requirements of the Securities Exchange Act of 1934, this report has been signed below on the 22nd day of November, 2017 by the following persons on behalf of the registrant and in the capacities indicated.

<u>Name</u>	<u>Capacity</u>
<u>/s/ VINCENT A. FORLENZA</u> Vincent A. Forlenza	Chairman and Chief Executive Officer (Principal Executive Officer)
<u>/s/ CHRISTOPHER R. REIDY</u> Christopher R. Reidy	Executive Vice President, Chief Financial Officer and Chief Administrative Officer (Principal Financial Officer)
<u>/s/ JOHN GALLAGHER</u> John Gallagher	Senior Vice President, Corporate Finance, Controller and Treasurer (Principal Accounting Officer)
<u>Basil L. Anderson*</u>	Director
<u>Catherine M. Burzik*</u>	Director
<u>R. Andrew Eckert*</u>	Director
<u>Claire M. Fraser*</u>	Director
<u>Christopher Jones*</u>	Director
<u>Marshall O. Larsen*</u>	Director
<u>Gary A. Mecklenburg*</u>	Director

<u>Name</u>	<u>Capacity</u>
_____ James F. Orr*	Director
_____ Willard J. Overlock, Jr.*	Director
_____ Claire Pomeroy*	Director
_____ Rebecca W. Rimel*	Director
_____ Bertram L. Scott*	Director

*By: _____ /s/ GARY DEFAZIO
 Gary DeFazio
 Attorney-in-fact

EXHIBIT INDEX

<u>Exhibit Number</u>	<u>Description</u>	<u>Method of Filing</u>
2(a)	Agreement and Plan of Merger, dated as of April 23, 2017, among C.R. Bard, Inc., Becton, Dickinson and Company and Lambda Corp.	Incorporated by reference to Exhibit 2.1 to the registrant's Current Report on Form 8-K dated April 24, 2017.
2(b)	Amendment No. 1, dated July 28, 2017, to the Agreement and Plan of Merger, dated as of April 23, 2017, among C.R. Bard, Inc., Becton, Dickinson and Company and Lambda Corp.	Incorporated by reference to Exhibit 2.1 to the registrant's Current Report on Form 8-K dated July 28, 2017.
3(a)(i)	Restated Certificate of Incorporation, dated as of January 29, 2013.	Incorporated by reference to Exhibit 3(a) to the registrant's Quarterly Report on Form 10-Q for the period ended March 31, 2013.
3(a)(ii)	Certificate of Amendment of the Restated Certificate of Incorporation, filed with the State of New Jersey Department of Treasury and effective May 15, 2017.	Incorporated by reference to Exhibit 4.1 to the registrant's registration statement on Form 8-A filed on May 16, 2017.
3(b)	By-Laws, as amended and restated as of April 23, 2017.	Incorporated by reference to Exhibit 3.1 to the registrant's Current Report on Form 8-K dated April 24, 2017.
4(a)	Indenture, dated as of March 1, 1997, between the registrant and The Bank of New York Mellon Trust Company, N.A. (as successor to JPMorgan Chase Bank)	Incorporated by reference to Exhibit 4(a) to Form 8-K filed by the registrant on July 31, 1997
4(b)	Form of 7% Debentures due August 1, 2027.	Incorporated by reference to Exhibit 4(d) of the registrant's Current Report on Form 8-K filed on July 31, 1997.
4(c)	Form of 6.70% Debentures due August 1, 2028.	Incorporated by reference to Exhibit 4(d) of the registrant's Current Report on Form 8-K filed on July 29, 1999.
4(d)	Form of 4.90% Notes due April 15, 2018.	Incorporated by reference to Exhibit 4(i) of the registrant's Annual Report on form 10-K for the fiscal year ended September 30, 2016.
4(e)	Form of 2.133% Notes due June 6, 2019.	Incorporated by reference to Exhibit 4.1 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(f)	Form of 2.675% Notes due December 15, 2019.	Incorporated by reference to Exhibit 4.3 of the registrant's Current Report on Form 8-K filed on December 15, 2014.
4(g)	Form of 2.404% Notes due June 5, 2020.	Incorporated by reference to Exhibit 4.2 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(h)	Form of 3.25% Notes due November 12, 2020.	Incorporated by reference to Exhibit 4.1 of the registrant's Current Report on Form 8-K filed on November 12, 2010.
4(i)	Form of 3.125% Notes due November 8, 2021.	Incorporated by reference to Exhibit 4.2 of the registrant's Current Report on Form 8-K filed on November 8, 2011.
4(j)	Form of 2.894% Notes due June 6, 2022.	Incorporated by reference to Exhibit 4.3 of the registrant's Current Report on Form 8-K filed on June 5, 2017.

<u>Exhibit Number</u>	<u>Description</u>	<u>Method of Filing</u>
4(k)	Form of 3.363% Notes due June 6, 2024.	Incorporated by reference to Exhibit 4.5 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(l)	Form of 3.734% Notes due December 15, 2024.	Incorporated by reference to Exhibit 4.4 of the registrant's Current Report on Form 8-K filed on December 15, 2014.
4(m)	Form of 3.700% Notes due June 6, 2027.	Incorporated by reference to Exhibit 4.6 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(n)	Form of 6.00% Notes due May 15, 2039.	Incorporated by reference to Exhibit 4.2 of the registrant's Current Report on Form 8-K filed on May 13, 2009.
4(o)	Form of 5.00% Notes due November 12, 2040.	Incorporated by reference to Exhibit 4.2 of the registrant's Current Report on Form 8-K filed on November 12, 2010.
4(p)	Form of 4.685% Notes due December 15, 2044.	Incorporated by reference to Exhibit 4.5 of the registrant's Current Report on Form 8-K filed on December 15, 2014.
4(q)	Form of 4.669% Notes due June 6, 2047.	Incorporated by reference to Exhibit 4.7 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(r)	Form of Floating Rate Notes due June 6, 2022.	Incorporated by reference to Exhibit 4.4 of the registrant's Current Report on Form 8-K filed on June 5, 2017.
4(s)	Form of 3.300% Senior Notes due March 1, 2023.	Incorporated by reference to Exhibit 4.4 of the registrant's Current Report on Form 8-K filed on April 29, 2015.
4(t)	Form of 3.875% Senior Notes due May 15, 2024.	Incorporated by reference to Exhibit 4.5 of the registrant's Current Report on Form 8-K filed on April 29, 2015.
4(u)	Form of 4.875% Senior Notes due May 15, 2044.	Incorporated by reference to Exhibit 4.6 of the registrant's Current Report on Form 8-K filed on April 29, 2015.
4(v)	Form of Certificate for the 6.125% Mandatory Convertible Preferred Stock, Series A.	Incorporated by reference to Exhibit 4.2 to the registrant's registration statement on Form 8-A filed on May 16, 2017.
4(w)	Deposit Agreement, dated as of May 16, 2017, among Becton, Dickinson and Company and Computershare Inc. and Computershare Trust Company, N.A., acting jointly as depository and Computershare Trust company, N.A., acting as Registrar and Transfer Agent, on behalf of the holders from time to time of the depository receipts described therein.	Incorporated by reference to Exhibit 4.3 to the registrant's registration statement on Form 8-A filed on May 16, 2017.
4(x)	Form of Depositary Receipt for the Depositary Shares.	Incorporated by reference to Exhibit 4.4 to the registrant's registration statement on Form 8-A filed on May 16, 2017.
10(a)(i)	Form of Employment Agreement with executive officers relating to employment following a change of control of the registrant (with tax reimbursement provisions).*	Incorporated by reference to Exhibit 10(a) to the registrant's Quarterly Report on Form 10-Q for the period ended December 31, 2008.

<u>Exhibit Number</u>	<u>Description</u>	<u>Method of Filing</u>
10(a)(ii)	Form of Employment Agreement with executive officers relating to employment following a change of control of the registrant (without tax reimbursement provisions).*	Incorporated by reference to Exhibit 10(a)(ii) to the registrant's Annual Report on Form 10-K for the fiscal year ended September 30, 2013.
10(b)	Stock Award Plan, as amended and restated as of January 31, 2006.*	Incorporated by reference to Exhibit 10(a) to the registrant's Quarterly Report on Form 10-Q for the period ended December 31, 2005.
10(c)	Performance Incentive Plan, as amended and restated January 24, 2017.*	Incorporated by reference to Exhibit 10.1 to the registrant's Quarterly Report on Form 10-Q for the period ended March 31, 2017.
10(d)	Deferred Compensation and Retirement Benefit Restoration Plan, as amended and restated as of September 27, 2016.*	Incorporated by reference to Exhibit 10(d) to the registrant's Annual Report on Form 10-K for the fiscal year ended September 30, 2016.
10(e)	1996 Directors' Deferral Plan, as amended and restated as of November 25, 2014.*	Incorporated by reference to Exhibit 10.2 to the registrant's Current Report on Form 8-K dated December 2, 2014.
10(f)	Amended and Restated Aircraft Time Sharing Agreement between Becton, Dickinson and Company and Vincent A. Forlenza dated as of March 21, 2012.*	Incorporated by reference to Exhibit 10.1 to the registrant's Current Report on Form 8-K dated March 27, 2012.
10(g)(i)	2004 Employee and Director Equity-Based Compensation Plan, as amended and restated as of January 26, 2016.*	Incorporated by reference to Exhibit 10 to the registrant's Current Report on Form 8-K dated January 29, 2016.
10(g)(ii)	Terms of Awards under 2004 Employee and Director Equity-Based Compensation Plan and Stock Award Plan.*	Incorporated by reference to Exhibit 10(g)(ii) to the registrant's Annual Report on Form 10-K for the fiscal year ended September 30, 2016.
10(h)	Five-Year Credit Agreement, dated January 29, 2016 among the registrant and the banks named therein (term has been extended to January 24, 2022).	Incorporated by reference to Exhibit 10 to the registrant's Current Report on Form 8-K dated February 4, 2016.
10(i)	364-Day Term Loan Agreement, dated December 19, 2014, by and among Becton, Dickinson and Company, as borrower, Goldman Sachs Bank USA, as administrative agent, and the lenders party thereto.	Incorporated by reference to Exhibit 10.1 to the registrant's Current Report on Form 8-K filed December 14, 2014.
10(j)	Form of Commercial Paper Dealer Agreement.	Incorporated by reference to Exhibit 10.1 to the registrant's Current Report on Form 8-K filed on January 6, 2015.
10(k)	Tax Matters Agreement, dated August 31, 2009, by and between Cardinal Health, Inc. and CareFusion Corporation.	Incorporated by reference to Exhibit 10.3 to Cardinal Health, Inc.'s Current Report on Form 8-K filed on September 4, 2009.
10(l)	Letter of Understanding dated March 28, 2016 between Becton, Dickinson and Company and Alexandre Conroy.*	Incorporated by reference to Exhibit 10 to the registrant's Quarterly Report on Form 10-Q for the period ended December 31, 2016.
10(m)	Three-Year Term Loan Agreement, dated as of May 12, 2017, by and among Becton, Dickinson and Company, the lenders party thereto, and Citibank, N.A., as administrative agent.	Incorporated by reference to Exhibit 10.1 to the registrant's Current Report on Form 8-K filed May 10, 2017.

<u>Exhibit Number</u>	<u>Description</u>	<u>Method of Filing</u>
10(n)	Credit Agreement, dated as of May 12, 2017, by and among Becton, Dickinson and Company, the banks and issuers of letters of credit party thereto and Citibank, N.A., as administrative agent.	Incorporated by reference to Exhibit 10.2 to the registrant's Current Report on Form 8-K filed May 10, 2017.
21	Subsidiaries of the registrant.	Filed with this report
23	Consent of independent registered public accounting firm.	Filed with this report
24	Power of Attorney.	Filed with this report
31	Certifications of Chief Executive Officer and Chief Financial Officer, pursuant to SEC Rule 13(a)-14(a).	Filed with this report
32	Certifications of Chief Executive Officer and Chief Financial Officer, pursuant to Section 1350 of Chapter 63 of Title 18 of the U.S. Code.	Filed with this report
101	The following materials from this report, formatted in XBRL (Extensible Business Reporting Language): (i) the Consolidated Statements of Income, (ii) the Consolidated Statements of Comprehensive Income, (iii) the Consolidated Balance Sheets, (iv) the Consolidated Statements of Cash Flows, and (v) Notes to Consolidated Financial Statements.	Filed with this report

* Denotes a management contract or compensatory plan or arrangement.

Copies of any Exhibits not accompanying this Form 10-K are available at a charge of 10 cents per page by contacting: Investor Relations, Becton, Dickinson and Company, 1 Becton Drive, Franklin Lakes, New Jersey 07417-1880, Phone: 1-800-284-6845.

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Corporate Officers

Vincent A. Forlenza

Chairman and Chief Executive Officer

Pierre A. Boisier

Executive Vice President and Chief Quality Officer

James W. Borzi

Executive Vice President, Global Operations and Chief Supply Chain Officer

Gary M. Cohen

Executive Vice President, Global Health and President, BD Foundation

Alexandre Conroy

President, Medication and Procedural Solutions

Gary M. DeFazio

Senior Vice President, Corporate Secretary and Associate General Counsel

John E. Gallagher

Senior Vice President, Corporate Finance, Controller and Treasurer

Roland Goette

Executive Vice President and President, EMEA

David W. Highet

Senior Vice President, Chief Intellectual Property Counsel and Assistant Secretary

James Lim

Executive Vice President and President, Greater Asia

Alberto Mas

Executive Vice President and President, Life Sciences Segment

George J. Parr

Executive Vice President and Chief Marketing Officer

Linda J. Peters

Executive Vice President and Chief Regulatory Officer

Aaron Pettit

Senior Vice President and Chief Ethics and Compliance Officer

Richard Pierle

Executive Vice President and Chief Information Officer

Thomas E. Polen

President

Christopher R. Reidy

Executive Vice President, Chief Financial Officer and Chief Administrative Officer

Antoinette F. Segreto

Senior Vice President, Taxes

Nabil Shabshab

President, Diabetes Care and Digital Health

Ellen R. Strahlman

Executive Vice President, Research and Development and Chief Medical Officer

Linda M. Tharby

Executive Vice President and Chief Human Resource Officer

Board of Directors

Basil L. Anderson^{1,2,4}

Retired Vice Chairman—Staples, Inc.

Catherine M. Burzik^{1,5}

Former President and Chief Executive Officer — Kinetic Concepts, Inc.

R. Andrew Eckert^{3,5}

President and Chief Executive Officer—Acelyty L.P. Inc.

Vincent A. Forlenza⁴

Chairman and Chief Executive Officer

Claire M. Fraser, Ph.D.^{3,5}

Director—Institute for Genome Sciences, University of Maryland School of Medicine

Christopher Jones^{1,4,5}

Retired Chief Executive Officer—JWT Worldwide

Marshall O. Larsen^{2,3,4}

Retired Chairman, President and Chief Executive Officer—Goodrich Corporation

Gary A. Mecklenburg^{2,3}

Retired President and Chief Executive Officer—Northwestern Memorial HealthCare

James F. Orr^{2,3}

Retired Chairman and Chief Executive Officer—Convergys Corporation

Willard J. Overlock, Jr.^{1,4,5}

Retired Partner—Goldman, Sachs & Company

Claire Pomeroy, M.D.^{3,5}

President—The Albert and Mary Lasker Foundation

Rebecca W. Rimel^{1,5}

President and Chief Executive Officer—The Pew Charitable Trusts

Bertram L. Scott^{1,2}

Senior Vice President of Population Health—Novant Health

Committees appointed by the Board of Directors

1. Audit Committee
2. Compensation and Management Development Committee
3. Corporate Governance and Nominating Committee
4. Executive Committee
5. Science, Marketing, Innovation and Technology Committee

*BD would like to thank **Basil Anderson** and **James Orr** for their years of dedicated service as they retire from the Board of Directors in 2018. For more than a decade, their leadership has helped shape BD into the global healthcare leader that we know today.*

Corporate information

Annual meeting

Tuesday, January 23, 2018 at 1:00 p.m.
Four Seasons Hotel New York
57 East 57th Street
New York, NY 10022

This annual report is not a solicitation of proxies.

Transfer agent and registrar

Computershare Trust Company, N.A.

By regular mail

P.O. Box 505000
Louisville, KY 40233-5000

By overnight mail

462 South 4th Street, Suite 1600
Louisville, KY 40202
Toll free: 877.498.8861
Toll: 781.575.2879
www.computershare.com

Direct stock purchase plan

The Direct Stock Purchase Plan established through Computershare Trust Company, N.A., enhances the services provided to existing shareholders and facilitates initial investments in BD shares. Plan documentation and additional information may be obtained by calling Computershare Trust Company, N.A., at 877.498.8861, or by accessing the "Buy stock direct" feature located within the Investor Centre of Computershare's website at www.computershare.com.

NYSE Symbol: BDX

Independent auditors

Ernst & Young LLP

5 Times Square
New York, NY 10036-6530
Phone: 212.773.3000
www.ey.com

Shareholder information

As of November 30, 2017, BD had 13,044 shareholders of record. The BD Statement of Corporate Governance Principles, the BD Code of Conduct, the charters of the BD Committees of the Board of Directors, BD reports and statements filed with or furnished to the Securities and Exchange Commission and other information are posted on the BD website at bd.com/investors.

Shareholders may receive, without charge, printed copies of these documents, including the BD 2017 Annual Report on Form 10-K, including the financial statements and related schedules, by contacting:

Investor relations

BD

1 Becton Drive
Franklin Lakes, NJ 07417-1880
Phone: 800.284.6845

bd.com

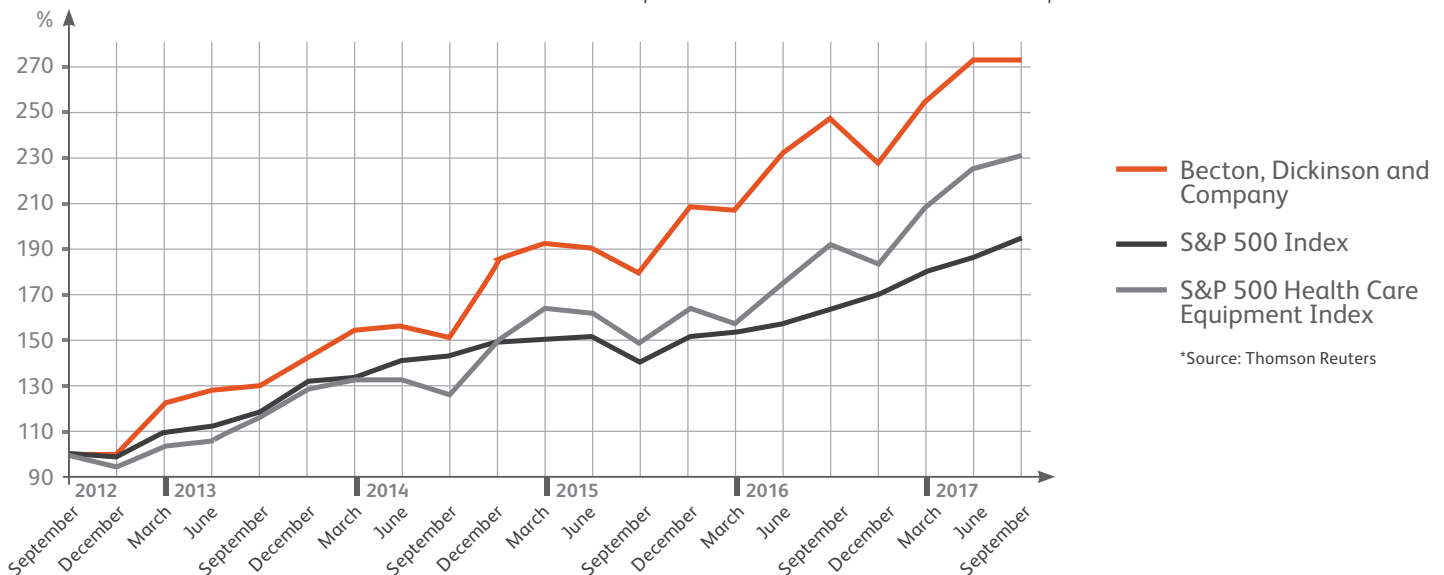
Comparison of five-year cumulative total return

1. Becton, Dickinson and Company
2. The S&P 500 Index
3. The S&P 500 Health Care Equipment Index*

The graph below presents a comparison of cumulative total return to shareholders for the five-year period ended September 30, 2017, for BD, the S&P 500 Index and the S&P 500 Health Care Equipment Index.

Cumulative total return to shareholders is measured by dividing total dividends (assuming dividend reinvestment) plus per share price change for the period by the share price at the beginning of the measurement period. The BD cumulative

shareholder return is based on an investment of \$100 on September 30, 2012, and is compared to the cumulative total return of the S&P 500 Index and the S&P 500 Health Care Equipment Index over the same period with a like amount invested.



*Source: Thomson Reuters



Franklin Lakes, NJ, 07417, U.S.

bd.com

