

SIRONA DENTAL SYSTEMS, INC.

Form 10-K

November 21, 2014

UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 10-K

(Mark One)

Annual Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

For the fiscal year ended September 30, 2014

or

Transition Report pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

For the transition period from _____ to _____

Commission file number 000-22673

Sirona Dental Systems, Inc.

(Exact name of registrant as specified in its charter)

Delaware

(State or other jurisdiction of incorporation or organization)

11-3374812

(I.R.S. Employer Identification No.)

30-30 47th Avenue, Suite 500, Long Island City, New York

(Address of principal executive offices)

11101

(Zip Code)

(718) 482-2011

(Registrant's telephone number, including area code)

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 Web site, 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, or a smaller reporting company. See the definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act. (Check one):

Large accelerated filer Accelerated filer Non-accelerated filer Smaller reporting company

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act).

Yes No

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The aggregate market value of common stock held by non-affiliates of the registrant as of March 31, 2014, the last business day of the registrant's most recently completed second fiscal quarter, was approximately \$2,519,469,890. Such aggregate market value is computed by reference to the closing sale price of the Common Stock on such date.

As of November 17, 2014, the number of shares outstanding of the Registrant's Common Stock, par value \$.01 per share, was 55,371,850.

DOCUMENTS INCORPORATED BY REFERENCE

Portions of the registrant's definitive proxy statement for its 2014 annual meeting of stockholders to be filed pursuant to Regulation 14A not later than 120 days after the end of the fiscal year (September 30, 2014) are incorporated by reference into Part III of this report on Form 10-K.

FORWARD-LOOKING STATEMENTS

This Form 10-K Annual Report contains forward-looking statements that involve risks and uncertainties. All statements, other than statements of historical facts, included in this Annual Report regarding the Company, its financial position, products, business strategy, and plans and objectives of management of the Company for future operations, are forward-looking statements. When used in this Annual Report, words such as “anticipate,” “believe,” “estimate,” “expect,” “intend,” “objectives,” “plans”, and similar expressions, or the negatives thereof or variations thereon or comparable terminology as they relate to the Company, its products or its management, identify forward-looking statements. Such forward-looking statements are based on the beliefs of the Company's management, as well as assumptions made by and information currently available to the Company's management. Actual results could differ materially from those contemplated by the forward-looking statements as a result of various factors, including, but not limited to, those contained in “Management's Discussion and Analysis of Financial Condition and Results of Operations” in Item 7 of this Annual Report and the “Risk Factors” set forth in Item 1A of this Annual Report. All forward looking statements speak only as of the date of this Annual Report and are expressly qualified in their entirety by the cautionary statements included in this report. The Company undertakes no obligation to update or revise forward-looking statements which may be made to reflect events or circumstances that arise after the date made or to reflect the occurrence of unanticipated events other than required by law.

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PART I

ITEM 1. BUSINESS

OVERVIEW

Sirona Dental Systems, Inc. (“Sirona,” the “Company,” “we,” “us,” and “our” refer to Sirona Dental Systems, Inc. and its consolidated subsidiaries and their predecessors, except as otherwise indicated or unless context otherwise requires) is the leading global manufacturer of high-quality, technologically-advanced dental equipment, and is focused on developing, manufacturing, and marketing innovative solutions for dentists around the world. The Company is uniquely positioned to benefit from several trends in the global dental industry, such as technological innovation, the shift to digital imaging, favorable demographic trends, and growing patient focus on dental health and cosmetic appearance. The Company’s headquarters is in Long Island City, New York, and its largest facility is located in Bensheim, Germany.

Sirona has a long tradition of innovation in the dental industry. The Company introduced the first dental electric drill approximately 130 years ago, the first dental X-ray unit approximately 100 years ago, the first dental computer-aided design/computer-aided manufacturing (CAD/CAM) system 28 years ago, and numerous other significant innovations in dentistry. Sirona continues to make significant investments in research and development (“R&D”), and its track record of innovative and profitable new products continues today. Sirona has the broadest product portfolio in the industry and is capable of fully outfitting and integrating a dental practice.

The majority of our revenues derive from the manufacture and sale of dental equipment. In addition, we also provide sales and after-sales service support to dentists and distributors through our growing sales and service infrastructure.

Sirona manages its commercial operations on both a product and geographic basis and maintains four reporting segments: 1) *Dental CAD/CAM Systems*, 2) *Imaging Systems*, 3) *Treatment Centers*, and 4) *Instruments*. Products from each category are marketed in all geographical sales regions.

The Company's business has grown substantially in the past several years, driven by numerous high-tech product introductions, a continued expansion of its global sales and service infrastructure, strong relationships with key distribution partners, namely Patterson and Henry Schein, and an international dealer network.

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Segment and geographic breakouts of Sirona's revenues for the fiscal year ended September 30, 2014 were as follows:

HISTORY

Sirona dates back to the establishment of Reiniger, Gebbert & Schall, which introduced the first dental electrical drill in 1882. In 1925, the Company became part of Siemens & Halske Group and in 1934 launched the smallest x-ray in the world, enabling dental x-rays for the first time. In 1956, Siemens introduced Sirona as a brand for treatment centers, and in 1958 the group developed the first ball-bearing turbine for dental drills.

In 1997, funds advised by the financial sponsor, Permira, acquired the Sirona dental business from Siemens in a leveraged buy-out transaction. Following the transaction, Sirona substantially increased its international sales and intensified its focus on product innovation. In November 2003, Permira sold Sirona to the Scandinavian financial sponsor, EQT, and management in a leveraged buy-out transaction that closed in February 2004. In April 2005, funds managed by Madison Dearborn Partners, a private equity firm, and Sirona's management entered into an agreement to acquire Sirona in a leveraged buy-out transaction that closed in June 2005.

In September 2005, Schick Technologies, Inc. ("Schick") entered into an Exchange Agreement with Sirona Holdings Luxco S.C.A. ("Luxco") and Sirona Holding GmbH ("Sirona Holding") providing for the issuance of 36,972,480 shares of Schick common stock to Luxco in exchange for Luxco's entire economic interest in Sirona Holding, which consisted of all of the issued and outstanding share capital of Sirona Holding and the existing indebtedness of Sirona Holding owed to Luxco in the principal amount of Euro 151.0 million (\$182 million) plus accrued interest (the "Exchange"). In June 2006, the Exchange closed and Schick, a Delaware corporation formed in 1997, was renamed Sirona Dental Systems, Inc. Although Sirona Holding became a subsidiary of Schick upon the completion of the Exchange, Sirona Holding was deemed the acquiring corporation for accounting purposes because Luxco received a controlling ownership interest in the Company, Sirona Holding's designees constituted a majority of the members of the Company's board of directors and Sirona Holding's senior management represented a majority of the senior

management of the Company. In May 2011, Luxco sold all of its remaining shares of Sirona common stock pursuant to an underwritten follow-on public offering.

Our common stock is currently traded publicly on the NASDAQ Global Select Market under the trading symbol "SIRO".

INDUSTRY/PRODUCTS

Overview

The global dental market encompasses the diagnosis, treatment and prevention of disease and ailments of the teeth, gums and supporting bone. This market has enjoyed steady growth, driven by a number of factors, including an increased desire for aesthetics, a demographic shift towards an aging population coupled with a desire to retain tooth structure later in life, growth in disposable income, a desire for more convenience on the part of both dentists and patients, a shift towards private pay, a greater need for dental preventative care.

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The global dental market has benefited from technological innovation, which increase productivity for the dentist. This is particularly important in markets facing increased demand for dental services with little or no increase in the number of dentists servicing those markets. In addition, technological developments allow dentists to offer higher quality treatment to patients. We believe that the high-tech end of the dental market is growing at a faster pace than the overall dental market and that this trend will continue over time.

Recent technological advancements in the dental equipment industry include 3D radiography, digital radiography, CAD/CAM technology, and intra oral cameras.

Sirona serves the high-tech dental equipment and technology market for dental practitioners and laboratories. We are the only manufacturing company that can fully outfit a dental practitioner's office with dental equipment, including treatment centers, imaging systems, dental CAD/CAM systems, and instruments. Our products represent important investments by dental practitioners, and some of these products can have a life span of 10-20 years (shorter for instruments and software), depending on the nature and quality of the product.

Products

Our principal products are generally classified into the following segments: *Dental CAD/CAM Systems, Imaging Systems, Treatment Centers, and Instruments.*

A brief description of each of our segments follows. See Note 10 to our consolidated financial statements for revenues and gross profit by segment for each of the last three fiscal years, and assets by segment, at September 30, 2014 and 2013.

Segment Revenue Contribution	Year ended		
	September 30,		
(In percent)	2014	2013	2012
Dental CAD/CAM Systems	37 %	37 %	34 %
Imaging Systems	34 %	35 %	35 %
Treatment Centers	19 %	19 %	20 %
Instruments	10 %	9 %	11 %
Total	100 %	100 %	100 %

Dental CAD / CAM Systems

Dental CAD/CAM Systems address the worldwide market for dental restorations, which includes several types of restorations, such as inlays, onlays, veneers, crowns, bridges, copings and bridge frameworks made from ceramic, metal or composite blocks. The global market for dental restorations can be divided into two sub-segments: in-mouth fillings and out-of-mouth pre-shaped restorations. CAD/CAM-produced ceramic restorations represent a growing portion of the out-of-mouth restoration market and the number of out-of-mouth restorations prepared with CAD/CAM systems has increased substantially over the past few years. At the same time, the number of dental practitioners and dental laboratories using CAD/CAM technology has increased. Sirona estimates that as of the end of fiscal year 2014, the market penetration for in-office CAD/CAM systems had grown to approximately 15% in the United States and Germany.

Sirona pioneered the application of high-tech CAD/CAM techniques to the traditional lab-based restoration process with the commercialization of the CERamic REConstruction, or CEREC, method. Sirona's CEREC system is an in-office application that enables dentists to produce high quality restorations from ceramic material and insert them into the patient's mouth during a single appointment. CEREC has a number of advantages compared to the traditional out-of-mouth pre-shaped restoration method, as CEREC does not require a physical model, restorations can be created in the dentist's office and the procedure can be completed in a single visit.

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The CEREC system consists of an imaging unit and a milling unit. The imaging unit scans the damaged area, captures the image of the tooth or teeth requiring restoration and proposes the specifications for the restoration. The milling unit then mills the ceramic restoration to the required specifications based upon the captured image and the dentist's design specifications. The result is a biocompatible, non-metallic, natural-looking restoration made of durable, high-quality ceramic materials completed in a single treatment session. Independent studies indicate that CEREC ceramic restorations are as durable as gold and can replace conventional restoration materials for most procedures. In addition, CEREC restorations are aesthetically pleasing and have the benefit of a natural-looking appearance.

Sirona offers a service contract on its CEREC product, which includes software updates and upgrades and maintenance on software-related hardware.

In addition to CEREC, Sirona also offers CAD/CAM products for dental laboratories, including the inLab restoration fabrication system and the extra-oral inEos scanner. These products are designed to improve efficiency and reduce costs for the dental lab. The inLab system scans the models received from the dentists and then mills ceramic or composite block restorations, such as crown copings and bridge frameworks to the specifications of the captured image.

Summary Innovation Timeline

(last 10 years)

Dental CAD/CAM Systems

Year Product / Event

- 2003 Launch of CEREC 3D software, an important development that allowed the dentist to view the onscreen restoration area in three dimensions.
- 2004 Start of the central restoration service business in Germany. The central restoration service allows dental labs to scan a plaster model received from the dentist and then transmit the digital image directly to Sirona via the internet. A restoration is then created at Sirona's central manufacturing site, with the final product shipped directly back to the lab.
- 2005 Launch of the inEos scanner, which is a high speed extra-oral scanner which produces 3D digital images from a single tooth up to a jaw, directly from the plaster model.
- 2007 Launch of the MC XL next generation milling unit, which produces high quality, precisely fitted restorations in about half the time that the older CEREC milling units required.

Introduction of Sirona's Biogeneric software, which virtually automated the design portion of the CAD/CAM process for inlays and onlays.

Launch of the next generation inLab milling unit, the inLab MC XL. The inLab MC XL milling and grinding unit opens up a broad range of production options for the dental laboratory. Milling performance and precision have been greatly enhanced, and a switch from grinding to milling can be accomplished in just a few, simple steps.

2008 Expansion of the central restoration service business to the United States.

Expanded our CEREC offering with the introduction of Sirona Connect. Sirona Connect is a web-based service that facilitates the electronic transmission of digital impressions acquired with a CEREC acquisition unit to inLab laboratories. Laboratories can use the digital impression to create final restorations. This process eliminates the need to take physical impressions, resulting in increased accuracy, less reworking of restorations and productivity savings.

2009 Launch of a new CEREC camera, based on the Company's proprietary Bluecam technology, which was faster, more accurate, and improved the workflow for practitioners.

2010 Further enhancement Sirona's Biogeneric software with the introduction of Version 3.8, which has the ability to create crowns and bridges.

Introduction of the CEREC AC Connect stand-alone digital impression unit. CEREC AC Connect allows dental professionals to scan digital impressions and then send them to the inLab® dental laboratory of their choice.

Introduction of the successor inEos scanner, the inEos Blue. inEos Blue is based on the Bluecam technology, is easy to use, fast, precise, flexible, and its auto capture function allows for substantial time savings.

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2011 Introduction of CEREC 4.0 software, an entirely redesigned software that gives CEREC users enhanced capabilities and speeds up the restoration process. In addition, CEREC 4.0 enables dentists to design and manufacture multiple restorations simultaneously, further enhancing productivity and profitability.

2012 Launch of new CEREC Omnicam camera, which allows dentists to generate precise whole-arch scans in the shortest possible time. Three features of the CEREC Omnicam are particularly notable: video streaming, digitization of jaw structures in their natural color, and powderless scanning of tooth surfaces. This introduction further strengthens Sirona's leadership position in the dental CAD/CAM market.

Introduction of inLab 4.0 software, which offers an extended spectrum of clinical applications. New design tools facilitate a customized and direct workflow. The completely revised platform provides a secure basis for integrating future applications.

2013 Launch of updated and expanded CEREC 4.2 software, further differentiated its CAD/CAM milling product line with the units MC X and MC XL Premium, and introduced the Apollo DI digital imaging system, all in March. These introductions expanded our portfolio and enable Sirona to offer "CAD/CAM for Everyone", an approach which seeks to address the various needs of the widest possible range of dentists.

Launch of enhanced inLab MC XL.

Introduction of the successor inEos scanner, the inEos X5. The 5-axis in Eos X5 is unrivaled in precision and has flexible handling, quick scanning times, and a comprehensive application spectrum for all digitization tasks.

Launch of inLab4.2 software, which introduced further applications, such as smart design with virtual articulation, smile design, and other features.

2014 Launch of CEREC 4.3 software, which enables carbide milling for optimal fabrication of zirconium oxide and polymer materials. Improved and simplified workflow for Omnicam. Virtual articulation is now calculated automatically. Integrated button to automatically upload the current case in Sirona Connect. CEREC 4.3 also allows a screw-retained implant workflow. Together with the introduction of additional TiBases and Scanpost, this leads to unsurpassed growth in implantology consumables.

Launch of precolored inCoris TZI C blocks that remove the need to use coloring liquids, thus simplifying the operating process with an improved result.

Imaging Systems

Imaging Systems comprise a broad range of systems for diagnostic imaging in the dental practice. Sirona has developed a comprehensive range of imaging systems for 2D or 3D, panoramic, and intra-oral applications that allow the dentist to provide a full range of diagnostics and treat the patient in a more efficient manner, resulting in safer, faster, and better dentistry.

Intra-oral x-ray systems use image-capture sensor devices, which are inserted into the mouth behind the diagnostic area, and typically take images of one or two teeth. Panoramic x-ray systems produce images of the entire jaw structure by means of an x-ray tube and an image capture device, which rotates around the head.

Summary Innovation Timeline

(last 10 years)

Imaging Systems

Year Product / Event

2004 Introduction of the next generation of 2D digital panoramic x-ray systems, the Orthophos XG line.

2006 Expansion of our imaging system product line to include Schick's intraoral sensor portfolio based on CMOS technology, as a result of the Exchange.

2007 Introduction of the GALILEOS Comfort 3-D imaging unit. Today, three-dimensional imaging is offering dentists advanced diagnostic and therapeutic options in the fields of surgery, implantology, prosthetics, orthodontics, and restorative dentistry. GALILEOS integrates these capabilities efficiently into dental practices.

2008 Launch of GALILEOS Compact, which is specifically tailored to meet the needs of the general practitioner.

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2009 Introduction of software that facilitates the integration of 3D X-ray volume (bone level data) with a CEREC AC CAD/CAM scan (surface level information). This software allows the practitioner to plan both the implant surgery and the prosthetic at the start of the implant treatment session. This integrated process reduces the number of treatment sessions, resulting in greater accuracy and superior implant alignment. With this new software, the dental practitioner can now place more focus on the desired aesthetic outcome throughout the entire treatment process.

2011 Introduction of the flagship Orthophos XG 3DReady, which provides dental practitioners with a wide variety of diagnostic possibilities and is upgradeable to a 3D unit. Other models of the family include the Orthophos XG 5 and the basic model Orthophos XG 3.

Launch of the Orthophos XG 3D imaging unit. This system gives the practitioner traditional 2D panoramic imaging capability and the ability to scan and view a large, 8x8 centimeter 3D field of view (a scan big enough to capture the entire jaw). Orthophos XG 3D is also available with cephalometric options, orthodontic, implant, and other specialty programs.

2012 Launch of the next generation of intraoral digital radiography – the Schick 33 sensor and image management system. Schick 33 is the most advanced sensor in dentistry, delivering an unparalleled combination of high-resolution images and dynamic image management.

2013 Launch of the GALILEOS Compact^{PLUS}, which is specifically tailored to meet the needs of the general practitioner, orthodontist, and oral surgeon.

2014 Launch of the SICAT function - the first software which visualizes real, patient-specific movement of the lower jaw within the 3D module of the GALILEOS.

Treatment Centers

Treatment Centers comprise a broad range of products from basic dentist chairs to sophisticated chair-based units with integrated diagnostic, hygiene, and ergonomic functionalities, as well as specialist centers used in preventative treatment and for training purposes. Sirona offers specifically configured products to meet the preferences of dentists within each region in which it operates. Sirona's treatment center configurations and system integration are designed to enhance productivity by creating a seamless workflow within the dental practice. Sirona's centers therefore allow the dentist to both improve productivity and increase patient satisfaction, significant factors in adding value to his or her practice.

Summary Innovation Timeline

(last 10 years)

Treatment Centers

Year Product / Event

- 2004 Commencement of the manufacture of basic treatment centers for developing markets via acquisition of a leading local manufacturer.
- 2008 Launch of the TENEO treatment center, which combines industry-leading technology with a timeless design that provides both patient and dentist with the ultimate in convenience and comfort.
- 2011 Introduction of the SINIUS treatment center, a comfort class treatment center, which enables the dentist to maximize time and flexibility of their practice. SINIUS is fully networked and is easily integrated into any dental practice.
- 2014 Launch of the INTEGO treatment center. INTEGO offers the design of the new generation of Sirona treatment centers with top quality and flexible configuration options at an attractive price. The fully-networked treatment center comes in two versions: INTEGO and INTEGO pro with extended functionality.

Instruments

Sirona offers a wide range of instruments, including handheld and power-operated handpieces for cavity preparation, endodontics, periodontology and prophylaxis, which are regularly updated and improved. The instruments are supplemented by multi-function tips, supply and suction hoses, as well as care and hygiene systems for instrument preparation. Sirona's instruments are often sold as packages in combination with treatment centers. Sirona's unique handpiece cleaning and sterilizing machine, the DAC Universal, is the only fully automatic system for instruments hygiene – with its unique features it has defined the standard of care for infection prevention in the dental office.

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Sirona intends to continue to strengthen the position of its Instruments segment as a diversified supplier of high-quality, reliable, user-friendly and cost-efficient dental instruments.

Summary Innovation Timeline

(last 10 years)

Instruments

Year Product / Event

- 2004 Expansion of our instruments product line into hygiene systems via acquisition of the Danish hygiene specialist, manufacturer of the popular DAC Universal.
- 2005 Launch of first generation SIROLaser - the smallest diode laser worldwide, redefining ergonomics in laser dentistry.
Launch of SIROEndo, a device for endodontic treatment with built-in APEX Location, seamlessly adaptable to the treatment center, and optimal integration into daily workflow.
- 2007 Launch of SIROPure - the first handpiece not requiring any lubrication over its entire product lifetime, keeps the cavity oil-free, eliminates maintenance errors, and lowers total cost of ownership.
- 2009 Launch of new diode laser, SIROLaser Advance / SIROLaser Xtend.
Launch of 3rd generation of DAC Universal, allowing the processing of wrapped instruments.
- 2011 Launch of SIROBoost high performance power turbine line with a high torque level, allowing faster, more efficient, and comfortable operation.
- 2012 Launch of the new handpiece program: T2 Line / T3 Line.
- 2013 Launch of new turbine generation (T1/T2/T3) and the T4 Line.
Launch of SIROInspect - fluorescence-based cavity detection device, whose safe and simple operation significantly reduces the risk of secondary cavities.
- 2014 Launch of new T3 Racer, with 30 watts the most powerful turbine on the market.

Manufacturing and Suppliers

Our main manufacturing and assembly activities are located in Bensheim, approximately 60 kilometers south of Frankfurt am Main, Germany. We also operate smaller manufacturing sites in New York, Italy, Denmark and China. All of our facilities are in good condition.

All of our manufacturing facilities have established and maintain a Quality Management System that is registered to ISO 9001:2000 and ISO 13485:2003. Our New York and Bensheim facilities also maintain a Device Establishment Registration with the United States Food and Drug Administration.

Manufacturing consists primarily of assembly, systems integration and testing. We generally outsource manufacturing of parts and components used in the assembly of our products but own the design and tools used by our key component suppliers. We do, however, manufacture most of the precision parts used for our instruments.

We purchase various components for our products from a number of outside suppliers. We currently have established relationships with approximately 1,400 suppliers, of which we view approximately 160 as “key suppliers.” Each supplier is selected according to stringent quality criteria, which are reviewed regularly. We do not believe we are dependent on one or a small group of suppliers and believe we could locate alternative suppliers if needed. For reasons of quality assurance or cost effectiveness, the Company relies on single sources for certain purchased components, e.g. sensors, which we use in our imaging segment. We work closely with our suppliers to help ensure continuity of supply while maintaining high quality and reliability. We have agreements in place and use a number of techniques, including security or consignment stock commitments, to address potential disruptions of the supply chain. We also own any custom tooling used in manufacturing these components. The Company has not experienced any significant difficulty in the past in obtaining the materials necessary to meet its production schedule. However, the need to replace one of our single source suppliers could cause a disruption in our ability to timely deliver certain of our products or increase costs.

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See Item 1A Risk Factors — “We are dependent upon a limited number of suppliers for critical components. If these suppliers delay or discontinue the manufacture of these components, we may experience delays in shipments, increased costs and cancellation of orders for our products.”

Sales and Marketing

Our sales and marketing efforts are directed through regional managers who oversee our sales professionals. These professionals work closely with our distribution partners to maximize the efficiency and productivity of their sales efforts. Our marketing initiatives are focused on highlighting Sirona’s leading role as a high-tech systems provider and industry innovator. In order to promote our brand and increase client loyalty, our distribution partners are supported through wide ranging advertising activities. In addition, we have been a key presenter at all major dental exhibitions, which are critical forums for raising brand awareness and new product introductions. Lastly, our product information is actively made available to business publications, dentists, journals, professional organizations and dental schools, our website (www.sirona.com), and social media offerings (Facebook, Twitter, Sirona Blog, etc.) are important interactive platforms for end-users as well as for distributors.

Distribution

Sirona distributes its products globally to dental practices, clinics and laboratories through an international network of more than 490 distributors and increasingly through our own sales and service infrastructure. See Note 10 to our consolidated financial statements for a description of our net sales and long-lived assets by geographic region for the last three fiscal years. Because distributors typically cover both dental equipment and consumables, they have regular contact with the dentist and are therefore optimally positioned to identify new equipment sale opportunities. Sirona’s primary distributors are Patterson Companies and Henry Schein, two of the world’s largest dental distributors. In the United States, Patterson is Sirona’s primary distributor. Outside of the United States, Henry Schein is the company’s largest distributor. Patterson Companies and Henry Schein accounted for 31% and 14%, respectively, of Sirona’s worldwide revenue for the fiscal year ended September 30, 2014. Sirona distributes elsewhere through a well-developed network of independent regional distributors. Sirona works closely with its distributors by training their technicians and sales representatives with respect to its products. With over 10,000 sales and service

professionals trained each year, Sirona seeks to ensure high standards of quality in after-sale service and the best marketing of its products. The success of Sirona's products is evidenced by their importance to its distribution partners, and in many cases are among their best-selling offerings. The Company continues to expand its sales and service infrastructure in selected countries around the world. These investments allow us to support our distributors' selling efforts and strengthen the Sirona brand in these key markets. These investments, and the subsequent expansion of our infrastructure, have enabled Sirona to grow revenues and profitability at a faster rate.

On April 27, 1998, Sirona and Patterson Companies entered into an exclusive distribution agreement (the "CAD/CAM Distribution Agreement") pursuant to which Patterson was appointed as the exclusive distributor of Sirona's CEREC CAD/CAM products within the United States and Canada. Under the terms of the CAD/CAM Distribution Agreement, Patterson's exclusivity was to terminate on September 30, 2007. On June 30, 2005, Sirona and Patterson entered into an amendment of the CAD/CAM Distribution Agreement which extended Patterson's exclusivity from October 1, 2007 through September 30, 2017. As consideration for the extension of its exclusivity, Patterson agreed to make a one-time payment to Sirona in the amount of \$100 million (the "Exclusivity Fee"). In July 2005, Patterson paid the Exclusivity Fee, in its entirety, to Sirona. The full amount of the Exclusivity Fee was recorded as deferred income and has been recognized on a straight-line basis since October 1, 2007. The extension did not modify or alter the underlying provisions of the companies' agreement through 2007, including the performance criteria necessary to maintain the exclusivity. The performance criteria are benchmark thresholds which afford Sirona the opportunity to abandon the exclusivity or to terminate the agreement with Patterson, but do not create minimum purchase obligations under a take-or-pay arrangement. The CAD/CAM Distribution Agreement was amended in May 2011 to revise the parameters for inLab sales in the United States and Canada.

In April 2000, Schick and Patterson entered into an exclusive distribution agreement (the "Schick Distribution Agreement") covering the United States and Canada; and as of May 1, 2000, Schick began marketing and selling its CDR dental products in the United States and Canada through Patterson.

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This contract was amended in July 2005, March 2007, and May 2010 and expired on December 31, 2012.

In May 2012, the Company and Patterson amended and restated the terms of their business relationship set forth in CAD/CAM Distribution Agreement and the Schick Distribution Agreement with respect to distribution of certain products throughout the United States and in October 2013 entered into new distribution agreements covering Canada. The amendment and restatement of both the CAD/CAM Distribution Agreement and Schick Distribution Agreement addressed issues related to pricing, termination and annual minimum purchase quotas, and provided growth targets which, if achieved, extend the companies' exclusivity period.

Sirona executes separate contracts with Henry Schein for each product group in each of the various jurisdictions in which Henry Schein distributes its products. The contracts governing most of the products distributed through Henry Schein are non-exclusive. Each of the contracts provides for minimum annual purchases, which are set annually. The contracts have terms of up to five years. Either party is entitled to terminate any of the contracts upon six months' notice but generally not before the third anniversary of the contract. Sirona may terminate a contract upon 30 days' notice in case of Henry Schein's default under the terms of the contract.

Competition

Competition in the global dental market is fragmented by both geography and products. We compete with a variety of companies, including large international companies as well as smaller companies that compete regionally or on a narrower product line. Sirona competes on the basis of its comprehensive and innovative product line and its global distribution network.

Research and Development

Sirona commits significant resources to research and development, with a particular focus on developing products that offer new diagnostic and treatment options, while increasing comfort for both users and patients and streamlining process efficiency. Research and development statistics for the last three fiscal years are as follows:

Research and Development (In millions, except otherwise noted)	Year ended September 30,		
	2014	2013	2012
Research and development expenses	\$64.6	\$60.2	\$52.6
Research and development expenses (% of revenue)	5.5 %	5.5 %	5.4 %
Number of R&D professionals employed globally	320	286	268

Sirona also cooperates in its research efforts with partners in research facilities and dental practices around the world. In fiscal year 2011, Sirona opened the Center of Innovation in Bensheim, Germany. The Center underscores Sirona's ongoing commitment to innovation in dentistry. Housing the majority of research and development professionals under one roof will ensure the Company maximum collaboration, creativity, technological advancement, and idea sharing.

Patents, Trade Secrets and Proprietary Rights

We seek to protect our intellectual property through a combination of patent, trademark and trade secret protection. We believe that our future success will depend in part on our ability to obtain and enforce patents for our products and processes, preserve our trade secrets and operate without infringing the proprietary rights of others.

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Patents

We have an active corporate patent program, the goal of which is to secure patent protection for our technology. Sirona owns and/or maintains approximately 860 patents and patent applications throughout the world. The patents expire at various dates through 2030. We also license or sublicense some of the technology used in our products from third parties.

Trademarks

We generally attempt to build brand awareness of our products through the use of trademark registrations. “Sirona,” “CEREC,” “Orthophos,” “Heliodont,” “inLab,” “CDR,” and “Schick” are some of our key registered trademarks. In addition, we have common law trademark rights in several other names we use commercially in connection with our products.

Trade Secrets

In addition to patent protection, we own trade secrets and proprietary know-how, which we seek to protect through agreements with employees and other appropriate individuals. These agreements generally allow assignment of confidential information developed by or made known to the individual by the Company during the course of the individual’s relationship with the Company as confidential and not to be disclosed to third parties, except in specific limited circumstances. The agreements also generally assign to the Company all inventions conceived by the individual in the course of rendering services to the Company. However, there can be no assurance that the Company will be successful in enforcing this policy in each case, that the Company would have adequate remedies available for any breach or that the Company’s trade secrets will not otherwise become known to, or independently developed by, its competitors.

Regulation

Medical Devices

Most of our products require certain forms of regulatory clearance, including, but not limited to, marketing clearance by the United States Food and Drug Administration (the “FDA”) in accordance with the Federal Food, Drug and Cosmetic Act, as amended (the “FD&C Act”) and by our Notified Body in accordance with the European Union’s Medical Device Directive 93/42/EEC (“MDD”).

The FDA and MDD review process typically requires extended proceedings pertaining to product safety and efficacy. We believe that our future success will depend to a large degree upon commercial sales of improved versions of our current products and sales of new products; we will not be able to market such products in the U.S. or in the European Union without FDA or MDD clearance, respectively. There can be no assurance that any products developed by us in the future will be granted clearance by applicable regulatory authorities or that additional regulations will not be adopted or current regulations amended in such a manner as to adversely affect us.

Pursuant to the FD&C Act, the FDA regulates the introduction, manufacture, advertising, labeling, packaging, marketing and distribution of, and record-keeping for, dental devices. The FDA classifies medical devices intended for human use into three classes: Class I, Class II, and Class III. The Company’s products are classified by the FDA into Class I or II that renders them subject only to general controls that apply to all medical devices, in particular regulations regarding alteration, misbranding, notification, record-keeping and good manufacturing practices.

The FD&C Act further provides that, unless exempted by regulation, medical devices may not be commercially distributed in the U.S. unless they have been cleared by the FDA. There are two review procedures by which medical devices can receive such clearance. Some products may qualify for clearance under a Section 510(k) procedure, in which the manufacturer submits to the FDA a pre-market notification that it intends to begin marketing the product, and shows that the product is substantially equivalent to another legally marketed product (i.e., that it has the same intended use and that it is as safe and effective as a legally marketed device, and does not raise different questions of safety and effectiveness than does a legally marketed device). Certain Class I devices are exempt from the 510(k) pre-market notification requirement and manufacturers of such products may proceed to market without any submission to the FDA. In some cases, the 510(k) notification must include data from human clinical studies.

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Marketing in the U.S. may commence once the FDA issues a clearance letter finding such substantial equivalence. According to FDA regulations, the agency has 90 days in which to respond to a Class I or II 510(k) notification. There can be no assurance, however, that the FDA will provide a timely response, or that it will reach a finding of substantial equivalence.

If a product does not qualify for the 510(k) procedure (either because it is not substantially equivalent to a legally marketed device or because it is a Class III device), the FDA must approve a Pre-Market Approval (“PMA”) application before marketing can begin. PMA applications must demonstrate, among other things, that the medical device is safe and effective. A PMA application is typically a complex submission that includes the results of clinical studies. Preparation of such an application is a detailed and time-consuming process. Once a PMA application has been submitted, the FDA’s review process may be lengthy and include requests for additional data. By statute and regulation, the FDA may take 180 days to review a PMA application, although such time may be extended. Furthermore, there can be no assurance that the FDA will approve a PMA application.

The products that we distribute in the European Union bear the “CE Mark,” a European Union symbol of compliance with the MDD. In order to market our products in the member countries of the European Union, it is necessary that those products conform to the requirements of the MDD. Our Bensheim facility which is engaged in the manufacturing of Class IIa and Class IIb medical devices as defined by the MDD is ISO 13485 certified. It is also necessary that our products comply with any revisions which may be made to these standards or the MDD.

Medical devices are subject to ongoing regulatory oversight by the FDA and a Notified Body. The FD&C Act requires that all medical device manufacturers and distributors register annually with the FDA and submit a list of those medical devices which they distribute commercially. The MDD requires that Class IIa devices or higher bear a CE mark with a Notified Body Number. The FD&C Act and the MDD also requires that all manufacturers of medical devices comply with labeling requirements and manufacture their products and maintain their documents in a prescribed manner with respect to manufacturing, testing, and quality control activities. The FDA’s Medical Device Reporting regulation and the MDD subject medical devices to post-market reporting requirements for death or serious injury, and for certain malfunctions that would be likely to cause or contribute to a death or serious injury if malfunction were to recur. In addition, the FDA and the MDD prohibit a device which has received marketing clearance from being marketed for applications for which marketing clearance has not been obtained. Furthermore, the FDA generally requires that medical devices not cleared for marketing in the U.S. receive FDA marketing clearance

before they are exported, unless an export certification has been granted. The FDA and the ISO Notified Bodies regularly inspect our registered and/or certified facilities.

Failure to comply with applicable regulatory requirements can, among other consequences, result in fines, injunctions, civil penalties, suspensions or loss of regulatory approvals, product recalls, seizure of products, operating restrictions and criminal prosecution. In addition, governmental regulations may be established that could prevent or delay regulatory clearance of our products. Delays in receipt of clearance, failure to receive clearance or the loss of previously received clearance would have a material adverse effect on our business, financial condition and results of operations.

Environmental, Health and Safety Matters

In addition to the laws and regulations discussed above, we are subject to government regulations applicable to all businesses, including, among others, regulations related to occupational health and safety, workers' benefits and environmental protection. The extent of government regulation that might result from any future legislation or administrative action cannot be accurately predicted. Failure to comply with regulatory requirements could have a material adverse effect on our business, financial condition and results of operations.

Employees

As of September 30, 2014, the Company had 3,327 employees. The Company believes that its relations with its employees are good. No Company employees are represented by labor unions or are subject to a collective bargaining agreement in the United States. Approximately 25% of our German employees are members of the IG Metall union. We have not experienced any work stoppages due to labor disputes.

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Executive Officers

See Part III, Item 10 of this 10-K Report for information about Executive Officers of the Company.

Available Information

Information about the Company's products and services, stockholder information, press releases, and filings with the Securities and Exchange Commission ("SEC") can be found on the Company's website at www.sirona.com. The information contained on our website is for informational purposes only and is not incorporated by reference into this report. The Company's Annual Reports on Form 10-K, Quarterly Reports on Form 10-Q, Current Reports on Form 8-K, and other SEC filings, and any amendments to such reports and filings, are available free of charge in the Investor Relations section of the Company's website as soon as reasonably practical after the Company's material is filed with, or furnished to, the SEC.

ITEM 1A. RISK FACTORS

INTRODUCTION

These risk factors may be important to understanding any statement in this Annual Report on Form 10-K or elsewhere. The following information should be read in conjunction with Management's Discussion and Analysis of Financial Condition and Results of Operations (MD&A), and the consolidated financial statements and related notes

incorporated by reference in this report.

Our businesses routinely encounter and address risks, some of which will cause our future results to be different - sometimes materially different - than we anticipate. Discussion about the material operational risks that our businesses encounter can be found in our MD&A, in the business descriptions in Item 1 of this report and in previous SEC filings. Below, we have described our present view of the material risks facing our business.

RISKS RELATED TO OUR BUSINESS

We must continue to develop new products and adapt to rapid technological change and evolving industry standards to remain competitive.

We are currently developing new products and enhancements to existing products. We cannot assure you that we will initiate, continue with and/or succeed in our efforts to develop or enhance such products. There can be no assurance that any new products will be developed by us, or if developed, will be approved by, or receive marketing clearance from, applicable domestic and/or international governmental or regulatory authorities.

The market for our products is characterized by rapid and significant technological change, evolving industry standards and new product introductions. Our products require significant planning, design, development and testing which requires significant capital commitments and investment by us. There can be no assurance that our products or proprietary technologies will not become noncompetitive or obsolete as a result of technological change, evolving industry standards or new product introductions or that we will be able to generate any economic return on our investment in product development. If our products or technologies become noncompetitive or obsolete, our business could be negatively affected.

If we cannot obtain or maintain approval from government agencies, we will not be able to sell our products.

We must obtain certain approvals by, and marketing clearances from, governmental authorities, including the FDA and similar health authorities in foreign countries to market and sell our products in those countries. These regulatory agencies regulate the marketing, manufacturing, labeling, packaging, advertising, sale and distribution of medical devices.

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The FDA enforces additional regulations regarding the safety of X-ray emitting devices. Our products are currently regulated by such authorities and certain of our new products will require approval by, or marketing clearance from, various governmental authorities, including the FDA. Various states also impose similar regulations.

The FDA review process typically requires extended proceedings pertaining to the safety and efficacy of new products. A 510(k) application is required in order to market a new or modified medical device. If specifically required by the FDA, a pre-market approval, or PMA, may be necessary. Such proceedings, which must be completed prior to marketing a new medical device, are potentially expensive and time consuming. They may delay or hinder a product's timely entry into the marketplace. Moreover, there can be no assurance that the review or approval process for these products by the FDA or any other applicable governmental authority will occur in a timely fashion, if at all, or that additional regulations will not be adopted or current regulations amended in such a manner as will adversely affect us. The FDA also oversees the content of advertising and marketing materials relating to medical devices which have received FDA clearance. Failure to comply with the FDA's advertising guidelines may result in the imposition of penalties.

We are also subject to other federal, state and local laws, regulations and recommendations relating to safe working conditions, laboratory and manufacturing practices. The extent of government regulation that might result from any future legislation or administrative action cannot be accurately predicted. Failure to comply with regulatory requirements could have a material adverse effect on our business.

Similar to the FDA review process, the EU review process typically requires extended proceedings pertaining to the safety and efficacy of new products. Such proceedings, which must be completed prior to marketing a new medical device, are potentially expensive and time consuming and may delay or prevent a product's entry into the marketplace.

Our profitability may be negatively impacted by adverse general macroeconomic conditions in the geographic markets in which we sell our products.

Our profitability depends in part on the varying economic and other conditions of the global dental market, which in turn is impacted by general macroeconomic conditions in the geographic markets in which we sell our products. Growth in the global dental market over the past few years has been driven by a number of factors, including a growth in disposable income, a shift towards private pay, a greater need for dental preventative care and an increased emphasis on aesthetics. Demand for our products would be negatively impacted by a decline in the economy in general, including interest rate and tax changes, that impact the financial strength of our customers, as well as by changes in the economy in general that reduce disposable income among dental consumers in the markets we sell our products, which would in turn reduce the demand for preventative and aesthetic dental services.

The ongoing disruptions in the overall world economy and financial markets could reduce disposable income among dental consumers and negatively affect the demand for dental services, which could be harmful to our financial position and results of operations. Furthermore, there can be no assurances that government responses to the disruptions in the financial markets will stabilize the markets or increase liquidity and the availability of credit for our customers. Difficult economic conditions may also result in a higher rate of losses on our accounts receivable. As a result, our business, results of operations or financial condition could be materially adversely affected.

We are dependent upon a limited number of distributors for a significant portion of our revenue, and loss of these key distributors could result in a loss of a significant amount of our revenue.

Historically, a substantial portion of our revenue has come from a limited number of distributors. For example, Patterson Dental Company, Inc. accounted for 31% of revenue for the fiscal year ended September 30, 2014. In addition, 14% of our revenue for the fiscal year ended September 30, 2014, was attributable to sales to Henry Schein, Inc. It is anticipated that Patterson and Henry Schein will continue to be the largest contributors to our revenue for the foreseeable future. There can be no assurance that Patterson and Henry Schein will purchase any specified minimum quantity of products from us or that they will continue to purchase any products at all. If Patterson or Henry Schein ceases to purchase a significant volume of products from us, it could have a material adverse effect on our results of operations and financial condition.

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Competition in the markets for our products is intense, and we may not be able to compete effectively.

Competition relating to our current products is intense and includes various companies, both within and outside of the United States. We anticipate that competition for our future products will also be intense and include various companies, both within and outside of the United States, Asia and Europe. Our competitors and potential competitors include large companies with substantially greater financial, sales and marketing, and technical resources, larger and more experienced research and development staffs, more extensive physical facilities and substantially greater experience in obtaining regulatory approvals and in marketing products than we have. In addition, we cannot assure you that our competitors are not currently developing, or will not attempt to develop, technologies and products that are more effective than those being developed by us or that would otherwise render our existing and new technology and products obsolete or noncompetitive. We may not be able to compete successfully and may lose market share to our competitors.

Our failure to obtain issued patents and, consequently, to protect our proprietary technology could hurt our competitive position.

Our success will depend in part on our ability to obtain and enforce claims in our patents directed to our products, technologies and processes, both in the United States and in other countries. Risks and uncertainties that we face with respect to our patents and patent applications include the following:

the pending patent applications that we have filed, or to which we have exclusive rights, may not result in issued patents or may take longer than we expect to result in issued patents;

- the allowed claims of any patents that issue may not provide meaningful protection;
- we may be unable to develop additional proprietary technologies that are patentable;

the patents licensed or issued to us may not provide a competitive advantage;

other companies may challenge patents licensed or issued to us;

disputes may arise regarding inventions and corresponding ownership rights in inventions and know-how resulting from the joint creation or use of intellectual property by us and our respective licensors; and

other companies may design around the technologies patented by us.

Our revenue and operating results are likely to fluctuate.

Our quarterly and annual operating results have varied in the past, and our operating results are likely to continue to fluctuate in the future. These variations result from a number of factors, many of which are substantially outside of our control, including:

- the timing of new product introductions by us and our competitors;
- timing of industry tradeshows, particularly the International Dental Show;
- changes in relationships with distributors;
- the timing of operational decisions by distributors and end users;
- developments in government reimbursement policies;
- changes in product mix;
- our ability to supply products to meet customer demand;
- fluctuations in manufacturing costs;
- tax incentives;

· currency fluctuations; and

- general economic conditions, as well as those specific to the healthcare industry and related industries.

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Our financial results may be adversely affected by fluctuations in foreign currency exchange rates.

We are exposed to currency exchange risk with respect to the U.S. Dollar in relation to the Euro, because a large portion of our revenue and expenses are denominated in Euros. In addition, we have an increasing portion of revenue and expenses denominated in other foreign currencies, e.g. Yen, Australian Dollar, Brazilian Real, and Yuan Renminbi. We monitor changes in our exposure to exchange rate risk. While we enter into hedging arrangements to protect our business against certain currency fluctuations, these hedging arrangements do not provide comprehensive protection, and our results of operations and prospects could be materially and adversely affected by foreign exchange fluctuations.

Our hedging and cash management transactions may expose us to loss or limit our potential gains.

As part of our risk management program, we use foreign currency exchange forward contracts. While intended to reduce the effects of exchange rate fluctuations, these transactions may limit our potential gains or expose us to loss. Should our counterparties to such transactions or the sponsors of the exchanges through which these transactions are offered fail to honor their obligations due to financial distress or otherwise, we would be exposed to potential losses or the inability to recover anticipated gains from these transactions.

We enter into foreign currency exchange forward contracts as economic hedges of trade commitments or anticipated commitments denominated in currencies other than the functional currency to mitigate the effects of changes in currency rates. Although we do not enter into these instruments for trading purposes or speculation, and although our management believes all of these instruments are economically effective as hedges of underlying physical transactions, these foreign exchange commitments are dependent on timely performance by our counterparties. Their failure to perform could result in our having to close these hedges without the anticipated underlying transaction and could result in losses if foreign currency exchange rates have changed.

We enter into interest rate swap agreements from time to time to manage some of our exposure to interest rate volatility. These swap agreements involve risks, such as the risk that counterparties may fail to honor their obligations under these arrangements. In addition, these arrangements may not be effective in reducing our exposure to changes in interest rates. If such events occur, our results of operations may be adversely affected.

Most of our cash deposited with banks is not insured and would be subject to the risk of bank failure. Our total liquidity also depends in part on the availability of funds under our Senior Facility Agreement. The failure of any bank in which we deposit our funds or that is part of our Senior Facility Agreement could reduce the amount of cash we have available for operations and additional investments in our business.

If we lose our key management personnel or are unable to attract and retain qualified personnel, it could adversely affect our results of operations or delay or hurt our research and product development efforts.

Our success is dependent, in part, upon our ability to hire and retain management, sales, technical, research and other personnel who are in high demand and are often subject to competing employment opportunities. It is possible that the loss of the services of one or a combination of our senior executives or key managers could have an adverse effect on our operations.

Work stoppages and other labor relations matters may make it substantially more difficult or expensive for us to produce our products, which could result in decreased sales or increased costs, either of which would negatively impact our financial condition and results of operations.

A significant part of our foreign employees are subject to collective bargaining agreements, and some of our employees are unionized; therefore, we are subject to the risk of work stoppages and other labor relations matters. While we have not experienced prolonged work stoppages in recent years and believe our relations with employees are satisfactory, any prolonged work stoppage or strike at any one of our principal facilities could have a negative impact on our business, financial condition, or results of operations.

We may experience difficulties managing our growth, which could adversely affect our results of operations.

It is expected that we will grow in certain areas of our operations as we develop and, assuming receipt of the necessary regulatory approvals, market our products. We will therefore need to recruit personnel, particularly sales and marketing personnel, and expand our capabilities, which may strain our managerial, operational, financial and other resources. To compete effectively and manage our growth, we must:

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·train, manage, motivate and retain a growing employee base;

·accurately forecast demand for, and revenue from, our product candidates; and

·expand existing operational, financial and management information systems to support our development and planned commercialization activities and the multiple locations of our offices.

Our failure to manage these challenges effectively could materially harm our business.

Since we operate in markets outside of the United States and Europe, we are subject to additional risks.

We anticipate that sales outside of the United States and Europe will continue to account for a significant percentage of our revenue. Such revenue is subject to a number of uncertainties, including, but not limited to, the following:

·economic and political instability;

·import or export licensing requirements;

·trade restrictions;

·longer payment cycles;

·unexpected changes in regulatory requirements and tariffs;

- fluctuations in currency exchange rates;
- potentially adverse tax consequences; and
- potentially weak protection of intellectual property rights.

These risks may impair our ability to generate revenue from our sales efforts. In addition, many countries outside of the United States and Europe have their own regulatory approval requirements for the sale of products. As a result, the introduction of new products into, and our continued sale of existing products in, these markets could be prevented, and/or costly and/or time-consuming, and we cannot assure you that we will be able to obtain the required regulatory approvals on a timely basis, if at all.

Our business is subject to extensive, complex, and changing laws, regulations, and orders that failure to comply with could subject us to civil or criminal penalties or other liabilities.

We are subject to extensive laws, regulations, and orders which are administered by various international, federal, and state governmental authorities, including, among others, the FDA, the Office of Foreign Assets Control of the United States Department of the Treasury (“OFAC”), the United States Federal Trade Commission, the United States Department of Justice, and other similar domestic and foreign authorities. These regulations include, but are not limited to, the U.S. Foreign Corrupt Practices Act and similar international anti-bribery laws, regulations concerning the supply of conflict minerals, various environmental regulations and regulations relating to trade, import and export controls and economic sanctions. Such laws, regulations, and orders may be complex and are subject to change.

Compliance with the numerous applicable existing and new laws, regulations and orders could require us to incur substantial regulatory compliance costs. Although the Company has implemented policies and procedures to comply with applicable laws, regulations and orders, there can be no assurance that governmental authorities will not raise compliance concerns or perform audits to confirm compliance with such laws, regulations, and orders. Failure to comply with applicable laws, regulations, or orders could result in a range of governmental enforcement actions, including fines or penalties, injunctions, and/or criminal or other civil proceedings. Any such actions could result in higher than anticipated costs or lower than anticipated revenue and could have a material adverse effect on the Company’s reputation, business, financial condition, and results of operations.

We may be exposed to liabilities under the Foreign Corrupt Practices Act, and any determination that we violated the Foreign Corrupt Practices Act could have a material adverse effect on our business.

To the extent that we operate outside the United States, we are subject to the Foreign Corrupt Practices Act (the “FCPA”) which generally prohibits U.S. companies and their intermediaries from bribing foreign officials for the purpose of obtaining or keeping business or otherwise obtaining favorable treatment. In particular, we may be held

liable for actions taken by our strategic or local partners even though such partners are foreign companies that are not subject to the FCPA.

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Any determination that we violated the FCPA could result in sanctions that could have a material adverse effect on our business.

New regulations related to conflict minerals could adversely impact our business.

The Dodd-Frank Wall Street Reform and Consumer Protection Act contains provisions designed to improve transparency and accountability concerning the supply of certain minerals, known as conflict minerals, originating from the Democratic Republic of Congo (DRC) and adjoining countries. As a result, in August 2012 the SEC adopted annual disclosure and reporting requirements for those companies who use conflict minerals mined from the DRC and adjoining countries in their products. These new requirements required due diligence efforts in fiscal 2013, with initial disclosure requirements beginning in May 2014. There will be additional costs associated with complying with these disclosure requirements, including for diligence to determine the sources of conflict minerals used in our products and other potential changes to products, processes or sources of supply as a consequence of such verification activities. The implementation of these rules could adversely affect the sourcing, supply, and pricing of materials used in our products. As there may be only a limited number of suppliers offering conflict-free minerals, we cannot be sure that we will be able to obtain necessary conflict minerals from such suppliers in sufficient quantities or at competitive prices. Also, we may face reputational challenges if we determine that certain of our products contain minerals not determined to be conflict free or if we are unable to sufficiently verify the origins for all conflict minerals used in our products through the procedures we may implement.

We may be a party to legal actions that are not covered by insurance.

We may be a party to a variety of legal actions, such as employment and employment discrimination-related suits, employee benefit claims, breach of contract actions, tort claims, stockholder suits, including securities fraud, governmental investigations and intellectual property related litigation. In addition, because of the nature of our business, we are subject to a variety of legal actions relating to our business operations. Although we have maintained insurance coverage for some of these potential liabilities, we cannot assure you that such insurance coverage will continue to be available or, if available, that it can be obtained in sufficient amounts or at reasonable cost or that it will be sufficient to cover any claims that may arise. Other potential liabilities may not be covered by insurance, insurers

may dispute coverage, or the amount of insurance may not be sufficient to cover the damages awarded. In addition, certain types of damages, such as punitive damages, may not be covered by insurance and/or insurance coverage for all or certain forms of liability may become unavailable or prohibitively expensive in the future.

We are dependent upon a limited number of suppliers for critical components. If these suppliers delay or discontinue the manufacture of these components, we may experience delays in shipments, increased costs and cancellation of orders for our products.

We rely on key suppliers for various critical components and procure certain components from outside sources which are sole suppliers. The availability and prices of these components may be subject to change due to interruptions in production, changing market conditions and other events. Any delays in delivery of or shortages in these components could interrupt and delay manufacturing of our products and result in the cancellation of orders for our products. In addition, these suppliers could discontinue the manufacture or supply of these components at any time. We may not be able to identify and integrate alternative sources of supply in a timely fashion or at all. Any transition to alternate suppliers may result in delays in shipment and increased expenses and may limit our ability to deliver products to our customers. If we are unable to develop reasonably-priced alternative sources in a timely manner, or if we encounter delays or other difficulties in the supply of such products and other materials from third parties, our business and results of operations may be harmed. In past years, semiconductors have been subject to significant price fluctuations.

While we have, in the past, attempted to mitigate the effects of such potential fluctuations, we cannot assure you that we will continue to do so or that we will be able to successfully mitigate the effect of future price increases on our results of operations and financial condition. See Item 1 Business – Manufacturing and Suppliers.

Our profitability could suffer if third parties infringe upon our proprietary technology.

Our profitability could suffer if third parties infringe upon our intellectual property rights or misappropriate our technologies and trademarks for their own businesses. To protect our rights to our intellectual property, we rely on a combination of patent and trademark law, trade secret protection, confidentiality agreements and contractual arrangements with our employees, strategic partners and others.

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We cannot assure you that any of our patents, any of the patents of which we are a licensee or any patents which may be issued to us or which we may license in the future, will provide us with a competitive advantage or afford us protection against infringement by others, or that the patents will not be successfully challenged or circumvented by third parties, including our competitors. The protective steps we have taken may be inadequate to deter misappropriation of our proprietary information. We may be unable to detect the unauthorized use of, or take appropriate steps to enforce, our intellectual property rights. Effective patent, trademark and trade secret protection may not be available in every country in which we will offer, or intend to offer, our products. Any failure to adequately protect our intellectual property could devalue our proprietary content and impair our ability to compete effectively. Further, defending our intellectual property rights could result in the expenditure of significant financial and managerial resources.

Our profitability may suffer if our products are found to infringe the intellectual property rights of others.

Litigation may be necessary to enforce our patents or to defend against any claims of infringement of patents owned by third parties that are asserted against us. In addition, we may have to participate in one or more interference proceedings declared by the United States Patent and Trademark Office, the European Patent Office or other foreign patent governing authorities, to determine the priority of inventions, which could result in substantial costs.

If we become involved in litigation or interference proceedings, we may incur substantial expense, and the proceedings may divert the attention of our technical and management personnel, even if we ultimately prevail. An adverse determination in proceedings of this type could subject us to significant liabilities, allow our competitors to market competitive products without obtaining a license from us, prohibit us from marketing our products or require us to seek licenses from third parties that may not be available on commercially reasonable terms, if at all. If we cannot obtain such licenses, we may be restricted or prevented from commercializing our products.

The enforcement, defense and prosecution of intellectual property rights, including the United States Patent and Trademark Office's, the European Patent Office's and other foreign patent offices' interference proceedings, and related legal and administrative proceedings in the United States and elsewhere, involve complex legal and factual questions. As a result, these proceedings are costly and time-consuming, and their outcome is uncertain. Litigation may be

necessary to:

- assert against others or defend us against claims of infringement;
- enforce patents owned by, or licensed to us from, another party;
- protect our trade secrets or know-how; or
- determine the enforceability, scope and validity of our proprietary rights or the proprietary rights of others.

Changes in the healthcare industry could adversely affect our business.

The healthcare industry has undergone, and is in the process of undergoing, significant changes driven by efforts to reduce costs. These changes include legislative healthcare reform, the reduction of spending budgets by government and private insurance programs, such as Medicare, Medicaid and corporate health insurance plans; trends toward managed care; consolidation of healthcare distribution companies; consolidation of healthcare manufacturers; collective purchasing arrangements and consolidation among office-based healthcare practitioners; and changes in reimbursements to customers. Some of these potential changes may cause a decrease in demand for and/or reduce the prices of our products. These changes could adversely affect our revenues and profitability. In addition, similar legislative efforts in the future could adversely impact our business.

The implementation of the Health Care Reform Law could adversely affect our business.

The Patient Protection and Affordable Care Act as amended by the Health Care and Education Reconciliation Act, each enacted in March 2010, generally known as the Health Care Reform Law, significantly expand health insurance coverage to uninsured Americans and changes the way health care is financed by both governmental and private payers. We expect expansion of access to health insurance to increase the demand for our products and services, but other provisions of the Health Care Reform Law could affect us adversely. Additionally, further federal and state proposals for health care reform are likely. We cannot predict what further reform proposals, if any, will be adopted, when they may be adopted, or what impact they may have on us.

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The Health Care Reform Law contains many provisions designed to generate the revenues necessary to fund the coverage expansions and to reduce costs of Medicare and Medicaid, including imposing a 2.3% excise tax on domestic sales of many medical devices by manufacturers and importers that began in 2013, which may adversely affect sales and cost of goods sold.

The implementation of the reporting and disclosure obligations of the Physician Payment Sunshine Act provisions of the Health Care Reform Law could adversely affect our business.

A Health Care Reform Law provision, generally referred to as the Physician Payment Sunshine Act or Open Payments Program, has imposed new reporting and disclosure requirements for drug and device manufacturers with regard to payments or other transfers of value made to certain practitioners (including physicians, dentists and teaching hospitals), and for such manufacturers and for group purchasing organizations, with regard to certain ownership interests held by physicians in the reporting entity. On February 1, 2013, the Centers for Medicare and Medicaid Services (“CMS”) released the final rule to implement the Physician Payment Sunshine Act. We expect to publish our first disclosure report in March 2015. As required under the Physician Payment Sunshine Act, CMS will publish information from these reports on a publicly available website, including amounts transferred and physician, dentist and teaching hospital identities.

The final rule implementing the Physician Payment Sunshine Act is complex, ambiguous, and broad in scope. CMS commentary on the final rule and more recent CMS communications indicate that wholesale drug and device distributors which take title to such products are to be treated as “applicable manufacturers” subject to full reporting requirements. In addition, certain of our subsidiaries manufacture devices. Accordingly, we are required to collect and report detailed information regarding certain financial relationships we have with dentists and teaching hospitals. It is difficult to predict how the new requirements may impact existing relationships among manufacturers, distributors, dentists and teaching hospitals. The Physician Payment Sunshine Act preempts similar state reporting laws, although we or our subsidiaries may be required to continue to report under certain of such state laws. While we expect to have substantially compliant programs and controls in place to comply with the Physician Payment Sunshine Act requirements, our compliance with the new final rule imposes additional costs on us.

If we fail to comply with laws and regulations relating to health care fraud, we could suffer penalties or be required to make significant changes to our operations, which could adversely affect our business.

We are subject to federal and state (and similar foreign) laws and regulations relating to health care fraud. Some of these laws, referred to as “false claims laws,” prohibit the submission or causing the submission of false or fraudulent claims for reimbursement to federal, state and other health care payers and programs. Other laws, referred to as “anti-kickback laws,” prohibit soliciting, offering, receiving or paying remuneration in order to induce the referral of a patient or ordering, purchasing, leasing or arranging for or recommending ordering, purchasing or leasing, of items or services that are paid for by federal, state and other health care payers and programs.

The government has expressed concerns about financial relationships between suppliers on the one hand and physicians and dentists on the other. As a result, we regularly review and revise our marketing practices as necessary to facilitate compliance. In addition, under the reporting and disclosure obligations of the Physician Payment Sunshine Act provisions of the Health Care Reform Law, the general public and government officials will be provided with new access to detailed information with regard to payments or other transfers of value to certain practitioners (including physicians, dentists and teaching hospitals) by applicable drug and device manufacturers subject to such reporting and disclosure obligations, which includes us. This information may lead to greater scrutiny, which may result in modifications to established practices and additional costs.

Failure to comply with health care fraud laws and regulations could result in significant civil and criminal penalties and costs, including the loss of licenses and the ability to participate in federal and state health care programs, and could have a material adverse impact on our business. Also, these laws may be interpreted or applied by a prosecutorial, regulatory or judicial authority in a manner that could require us to make changes in our operations or incur substantial defense and settlement expenses. Even unsuccessful challenges by regulatory authorities or private relators could result in reputational harm and the incurring of substantial costs. In addition, many of these laws are vague or indefinite and have not been interpreted by the courts, and have been subject to frequent modification and varied interpretation by prosecutorial, regulatory authorities, increasing compliance risks.

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While we believe that we are substantially compliant with the foregoing laws and regulations promulgated thereunder, and have adequate compliance programs and controls in place to ensure substantial compliance, we cannot predict whether changes in applicable law, or interpretation of laws, or changes in our services or marketing practices in response, could adversely affect our business.

If we fail to comply with laws and regulations relating to the confidentiality of sensitive personal information or standards in electronic health data transmissions, we could be required to make significant changes to our products, or incur penalties or other liabilities.

Certain of our businesses involve access to personal health, medical, financial and other information of individuals, and are accordingly directly or indirectly subject to numerous federal, state, local and foreign laws and regulations that protect the privacy and security of such information, and require, among other things, the implementation of various recordkeeping, operational, notice and other practices intended to safeguard that information, limit its use to allowed purposes, and notify individuals in the event of privacy and security breaches. Failure to comply with these laws can result in substantial penalties and other liabilities. As a result of the federal Health Information Technology for Economic and Clinical Health Act (“HITECH Act”), which was passed in 2009, some of our businesses that were previously only indirectly subject to federal Health Insurance Portability and Accountability Act of 1996 (“HIPAA”) privacy and security rules became directly subject to such rules because such businesses serve as “business associates” of HIPAA covered entities, such as health care providers. On January 17, 2013 the Office for Civil Rights of the Department of Health and Human Services released a final rule implementing the HITECH Act and making certain other changes to HIPAA privacy and security requirements. Compliance with the rule was required by September 23, 2013, and increases the requirements applicable to some of our businesses.

Product liability claims exposure could be significant.

We may face exposure to product liability claims and recalls for unforeseen reasons from consumers, distributors or others. We may experience material product liability losses in the future, and we may incur significant costs to defend

these claims. In addition, if any of our products are or are alleged to be defective; we may be required to participate in a recall involving those products. End-users of our products may look to us for contribution when faced with product recalls or product liability claims. Although we have maintained insurance coverage related to product liability claims, we cannot assure you that product liability insurance coverage will continue to be available or, if available, that it can be obtained in sufficient amounts or at reasonable cost or that it will be sufficient to cover any claims that may arise. We may not maintain any insurance relating to potential recalls of our products. A successful product liability claim brought against us in excess of available insurance coverage or a requirement to participate in any product recall could reduce our profits and/or impair our financial condition, and damage our reputation.

Product warranty claims exposure could be significant.

We generally warrant each of our products against defects in materials and workmanship for a period of one year from the date of shipment plus any extended warranty period purchased by the customer. The future costs associated with providing product warranties could be material. A successful warranty claim brought against us could reduce our profits and/or impair our financial condition, and damage our reputation.

Adverse publicity regarding the safety of our technology or products could negatively impact us.

Despite any favorable safety tests that may be completed with respect to our products, adverse publicity regarding application of X-ray products or other products being developed or marketed by others could negatively affect us. If other researchers' studies raise or substantiate concerns over the safety of our technology approach or product development efforts generally, our reputation could be harmed, which would adversely impact our business.

Inadequate levels of reimbursement from governmental or other third-party payers for procedures using our products may cause our revenue to decline.

Third-party payers, including government health administration authorities, private health care insurers and other organizations regulate the reimbursement of fees related to certain diagnostic procedures or medical treatments.

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Third-party payers are increasingly challenging the price and cost-effectiveness of medical products and services. While we cannot predict what effect the policies of government entities and other third-party payers will have on future sales of our products, there can be no assurance that such policies would not cause our revenue to decline.

We have developed and must continue to maintain internal controls.

Effective internal controls are necessary for us to provide assurance with respect to our financial reports and to effectively prevent fraud. If we cannot provide reasonable assurance with respect to our financial reports and effectively prevent fraud, our operating results could be harmed. The Sarbanes-Oxley Act of 2002 requires us to furnish a report by management on internal control over financial reporting, including managements' assessment of the effectiveness of such control. Internal control over financial reporting may not prevent or detect misstatements because of its certain limitations, including the possibility of human error, the circumvention or overriding of controls, or fraud. As a result, even effective internal controls may not provide reasonable assurances with respect to the preparation and presentation of financial statements. In addition, projections of any evaluation of effectiveness of internal control over financial reporting to future periods are subject to the risk that the control may become either obsolete or inadequate as a result of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate. If we fail to maintain adequate internal controls, including any failure to implement required new or improved controls, or if we experience difficulties in implementing new or revised controls, our business and operating results could be harmed and we could fail to meet our reporting obligations.

We may be required to record a significant charge to earnings if our goodwill or other intangible assets become impaired.

Our balance sheet includes goodwill and other identifiable intangible assets. If impairment of our goodwill or other identifiable intangible assets is determined, we may be required to record a significant charge to earnings in the period of such determination under U.S. generally accepted accounting principles (GAAP).

RISKS RELATED TO OUR COMMON STOCK

Certain provisions of our certificate of incorporation and bylaws and Delaware law could discourage, delay, or prevent a merger or acquisition at a premium price.

The provisions of our certificate of incorporation and bylaws may also deter, delay or prevent a third-party from acquiring us. These provisions include:

· limitations on the ability of stockholders to amend our charter documents, including stockholder supermajority voting requirements;

· the authority of the board of directors to adopt amendments to our bylaws without shareholder approval;

· the inability of stockholders to act by written consent or to call special meetings;

· a classified board of directors with staggered three-year terms;

· advance notice requirements for nominations for election to the board of directors and for stockholder proposals; and

· the authority of our board of directors to issue, without stockholder approval, up to 5,000,000 shares of preferred stock with such terms as the board of directors may determine and to issue additional shares of our common stock.

We are also subject to the protections of Section 203 of the Delaware General Corporation Law, which prevents us from engaging in a business combination with a person who acquires at least 15% of our common stock for a period of three years from the date such person acquired such common stock, unless board or stockholder approval were obtained.

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In addition, in the event of a “change of control” as defined in our senior facilities agreement, we may be required to, among other things, repay all of our obligations outstanding under the senior facilities agreement, with interest thereon, which could materially adversely impact the value of our common stock.

These provisions could have the effect of delaying, deferring or preventing a change in control of our company, discourage others from making tender offers for our shares, lower the market price of our stock or impede the ability of our stockholders to change our management, even if such changes would be beneficial to our stockholders.

The market price of our common stock may fluctuate significantly, and this may make it difficult for holders to resell our common stock when they want or at prices that they find attractive.

The price of our common stock on the NASDAQ Global Select Market constantly changes. We expect that the market price of our common stock will continue to fluctuate. The market price of our common stock may fluctuate as a result of a variety of factors, many of which are beyond our control. These factors include:

· changes in market conditions;

· quarterly variations in our operating results;

· operating results that vary from the expectations of management, securities analysts and investors;

· changes in expectations as to our future financial performance;

· announcements of strategic developments, significant contracts, acquisitions and other material events by us, our competitors, or our distribution partners;

- the operating and securities price performance of other companies that investors believe are comparable to us;
- future sales of our equity or equity-related securities;
- changes in the economy and the financial markets;
- departures of key personnel;
- changes in governmental regulations; and
- geopolitical conditions, such as acts or threats of terrorism or military conflicts.

In addition, in recent years, the stock market in general has experienced extreme price and volume fluctuations. This volatility has had a significant effect on the market price of securities issued by many companies for reasons often unrelated to their operating performance. These broad market fluctuations may adversely affect the market price of our common stock, regardless of our operating results.

Compliance with changing regulation of corporate governance and public disclosure will result in additional expenses and pose challenges for our management

Changing laws, regulations and standards relating to corporate governance and public disclosure, including the Dodd-Frank Wall Street Reform and Consumer Protection Act and the rules and regulations promulgated thereunder, the Sarbanes-Oxley Act and SEC regulations, have created uncertainty for public companies and significantly increased the costs and risks associated with accessing the U.S. public markets. Our management team will need to devote significant time and financial resources to comply with both existing and evolving standards for public companies, which will lead to increased general and administrative expenses and a diversion of management time and attention from revenue generating activities to compliance activities.

ITEM 1B. UNRESOLVED STAFF COMMENTS

None.

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ITEM 2. PROPERTIES

The Company leases its headquarters in Long Island City, New York. The lease expires in November 2017. The leased space houses executive offices and group functions including legal affairs and investor relations, sales and marketing, research and development laboratories and production and shipping facilities.

The Company has its largest facility in Bensheim, Germany. It is composed of a number of buildings housing the Company's primary manufacturing and assembly facility. It also houses executive offices, finance, sales, customer service and marketing, research and development laboratories, and shipping facilities. In fiscal year 2011, the Company expanded these facilities with inauguration of the Center of Innovation, which houses the research and development professionals in Germany under one roof. In fiscal year 2013, the Company once again invested in these facilities by significantly expanding and enhancing its Instruments manufacturing capacity. In addition, since September 2007, the Company leases space in Salzburg, Austria. The leased space houses executive offices and group functions including strategy, sales, finance, accounting, human resources, marketing, and legal affairs.

The Company also maintains manufacturing facilities in China, Italy and Denmark and certain sales and service offices worldwide.

The Company believes that its properties and facilities will be adequate for its needs for the foreseeable future and that, if such space proves to be inadequate, it will be able to procure additional or replacement space that will be adequate for its needs.

ITEM 3. LEGAL PROCEEDINGS

The Company is involved in various legal proceedings that are incidental to the conduct of the Company's business. The Company is not involved in any pending or threatened legal proceedings that the Company believes could reasonably be expected to have a material adverse effect on the financial condition or results of operations.

ITEM 4. MINE SAFETY DISCLOSURES

Not applicable.

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PART II

**ITEM MARKET FOR REGISTRANT'S COMMON EQUITY, RELATED STOCKHOLDER MATTERS
5. AND ISSUER PURCHASES OF EQUITY SECURITIES**

Our common stock is currently traded publicly on the NASDAQ Global Select Market. Our trading symbol is "SIRO".

QUARTERLY INFORMATION ON THE PRICE RANGE OF COMMON STOCK

The following table presents quarterly information on the price range of our common stock. This information indicates the high and low sale prices, as quoted on NASDAQ commencing October 1, 2012. These prices do not include retail markups, markdowns or commissions.

Quarterly Price Range of Common Stock

(In Dollars)	Range	
	High	Low
Fiscal Year Ended September 30, 2014		
First Quarter	\$73.94	\$66.04
Second Quarter	77.31	67.20
Third Quarter	82.84	72.11
Fourth Quarter	84.95	75.16
Fiscal Year Ended September 30, 2013		
First Quarter	\$64.57	\$53.26

Second Quarter	73.98	64.11
Third Quarter	75.81	62.48
Fourth Quarter	72.63	64.40

On November 17,, 2014, there were approximately 68 holders of record of the Company's common stock. However, the Company believes that the number of beneficial owners of its common stock is substantially higher.

Historically, Sirona has not paid any dividends to holders of its common stock. The Company may consider paying dividends in the future, but currently has no plans to do so. The payment of dividends is within the discretion of the Board of Directors and will depend upon the Company's earnings, its capital requirements, financial condition and other relevant factors.

For information relating to securities authorized for issuance under equity compensation plans, see Part III, Item 12.

ISSUER PURCHASES OF EQUITY SECURITIES

In May 2013, the Company's Board of Directors announced a stock repurchase program (the "2013 Program") to purchase up to an aggregate of \$100 million of its common stock in open market or privately-negotiated transactions effective through June 2016.

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Sirona Dental Systems Inc.	\$ 100.00	\$ 121.14	\$ 142.55	\$ 191.46	\$ 224.97	\$ 257.75
Russell 2000	100.00	113.35	109.35	144.24	187.59	194.96
Dow Jones US Medical Equipment	100.00	100.94	105.58	130.72	162.88	195.62

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The selected historical consolidated financial data of Sirona included below and elsewhere in this document are not necessarily indicative of future performance. This information is only a summary and should be read in conjunction with the sections entitled 'Management's Discussion and Analysis of Financial Condition and Results of Operations' and the consolidated financial statements contained elsewhere in this document.

Selected Income Statement Data	Year Ended				
	September 30,				
(In millions)	2014	2013	2012	2011	2010
REVENUE	\$1,171.1	\$1,101.5	\$979.4	\$913.9	\$770.3
GROSS PROFIT	641.7	591.4	524.0	483.7	399.0
OPERATING INCOME	238.1	212.8	185.8	160.9	128.1
INCOME BEFORE TAXES	230.4	197.5	178.3	159.5	115.2
NET INCOME	177.4	148.5	135.6	123.8	91.4
NET INCOME ATTRIBUTABLE TO SIRONA DENTAL SYSTEMS, INC.	\$175.7	\$146.7	\$133.8	\$121.8	\$90.0
INCOME PER SHARE					
(attributable to Sirona Dental Systems, Inc. common shareholders):					
Basic	\$3.18	\$2.67	\$2.41	\$2.19	\$1.63
Diluted	\$3.13	\$2.61	\$2.36	\$2.13	\$1.59

Selected Balance Sheet Data	As of				
	September 30,				
(In millions)	2014	2013	2012	2011	2010
CASH AND CASH EQUIVALENTS	\$382.8	\$241.7	\$151.1	\$345.9	\$251.8
WORKING CAPITAL ^{(1), (2)}	449.8	317.0	222.9	46.2	297.6
TOTAL ASSETS	1,811.0	1,738.4	1,494.7	1,726.1	1,592.9

NON-CURRENT LIABILITIES ⁽²⁾	312.5	334.3	315.9	255.0	625.2
TOTAL LIABILITIES	549.8	580.4	502.3	790.2	785.3
RETAINED EARNINGS	759.9	584.2	437.5	303.6	181.8
SIRONA DENTAL SYSTEMS, INC. SHAREHOLDERS' EQUITY	1,258.8	1,155.6	989.4	932.3	805.4
TOTAL SHAREHOLDERS' EQUITY	1,261.2	1,158.0	992.4	935.9	807.6
NET CASH (DEBT) ⁽³⁾	\$303.3	\$166.3	\$75.6	\$(22.5)	\$(119.0)

⁽¹⁾ Working capital is defined as current assets less current liabilities.

⁽²⁾ The significant decrease in working capital and non-current liabilities in fiscal year 2011 is due to the reclassification of the final tranche of the senior term loan due in November 2011 as current. The balance of these senior term loans was \$364.8 as of September 30, 2011.

⁽³⁾ Net cash (debt) is defined as cash and cash equivalents less short and long-term financial liabilities.

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

INTRODUCTION

Except as otherwise indicated or unless context otherwise requires, the terms "Sirona", the "Company", "we", "us", and "our" refer to Sirona Dental Systems, Inc. and its consolidated subsidiaries.

Management's Discussion and Analysis of Financial Condition and Results of Operations (MD&A) is intended to provide the reader of our financial statements with a narrative, from the perspective of our management, on our business, financial condition, results of operations, liquidity, and certain other factors that may affect our future results. Unless expressly stated otherwise, the comparisons presented in this MD&A refer to the same period in the prior year.

Our MD&A should be read in conjunction with the Consolidated Financial Statements included elsewhere in this report. Actual results and the timing of certain events may differ significantly from those projected in forward-looking statements due to a number of factors, including those set forth in "Operations Review" in this Item and elsewhere in this Report.

Certain revenue information is presented on a constant currency basis ("Constant Currency"). This information is a non-GAAP financial measure. Sirona supplementally presents this revenue information because it believes doing so facilitates a comparison of its operating results from period to period without regard to changes resulting solely from

fluctuations in currency rates. Because of the historical significance of the Euro to its operations, Sirona calculates Constant Currency revenue growth by comparing current-period revenues to prior-period revenues with both periods converted at the U.S. Dollar/Euro average foreign exchange rate for each month of the current period. Currently, our Constant Currency calculation does not adjust for fluctuations in any other currency, most importantly the Japanese Yen, Chinese Yuan Renminbi, Australian Dollar, and Brazilian Real.

All amounts in this section are reported in millions of U.S. Dollars (\$), except as otherwise disclosed.

EXECUTIVE OVERVIEW OF SIRONA'S BUSINESS AND PERFORMANCE

Business

Sirona is the leading global manufacturer of high-quality, technologically-advanced dental equipment, and is focused on developing, manufacturing, and marketing innovative systems and solutions for dentists around the world. The Company is uniquely positioned to benefit from several trends in the global dental industry, such as technological innovation, increased use of CAD/CAM systems in restorative dentistry, the shift to digital imaging, favorable demographic trends, and growing patient focus on dental health and cosmetic appearance. The Company's headquarters is in Long Island City, New York, and its largest facility is located in Bensheim, Germany.

Sirona has a long tradition of innovation in the dental industry. The Company introduced the first dental electric drill over 130 years ago, the first dental X-ray unit approximately 100 years ago, the first dental computer-aided design/computer-aided manufacturing (CAD/CAM) system 29 years ago, and numerous other significant innovations in dentistry. Sirona continues to make significant investments in R&D, and its track record of innovative and profitable new products continues today. Sirona has the broadest product portfolio in the industry and is capable of fully outfitting and integrating a dental practice.

Sirona manages its commercial operations on both a product and geographic basis and maintains four reporting segments: 1) *Dental CAD/CAM Systems*, 2) *Imaging Systems*, 3) *Treatment Centers*, and 4) *Instruments*. Products from each category are marketed in all geographical sales regions.

Our business has grown substantially in the past several years, driven by numerous high-tech product introductions and continued expansion of our global sales and service infrastructure. Our growth was enabled by continued development of strong relationships with key distribution partners, namely Patterson in the U.S. and Henry Schein in Europe.

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The U.S. market is the largest individual market for Sirona, followed by Germany. We have also pursued expansion in other international markets. Over the last five fiscal years, the Company's geographical revenue development was as follows:

U.S.: Revenues have been driven by innovative products, particularly in the CAD/CAM and Imaging segments.

International Markets: Sirona has been able to grow revenues in international markets by gaining market share with its innovative, best-in-class product solutions as well as through expansion of its local presence and distribution channels by establishing sales and service locations in countries such as Australia, Brazil, China, Italy, Japan, Russia, Slovakia, South Africa, South Korea, and Turkey. The expansion helped to increase market share but also contributed to higher SG&A expenses.

Sirona has been able to grow by often being first-to-market and establishing a strong distribution network as these countries' dental markets expanded. Additionally, increasing demand for best-in-class dental technology and growing middle class populations demanding better dental care in these markets have increased demand for Sirona's products.

Current Performance at a Glance

The following is a synopsis of our performance for the fiscal year ended September 30, 2014:

Revenue: For the fiscal year ended September 30, 2014, we reported revenue growth of 6.3%. On a Constant Currency basis, total revenues were up 3.9% over a strong fiscal year 2013, where revenues grew 11.7% Constant Currency. In both fiscal years, Constant Currency revenues were negatively impacted by exchange rate fluctuations other than the U.S. Dollar/Euro exchange rate. We continue to experience solid growth in demand for our CAD/CAM and Imaging products. Instruments was the strongest growing segment this year benefiting from our new state-of-the-art manufacturing facility in Bensheim, Germany, as well as strong demand for our hygiene product line. With the launch of our new Intego treatment center and various trade-up programs in the current fiscal year, Treatment Centers increased over a very strong prior year, which was positively impacted by the last-edition program for our popular M1+ unit. Geographically, growth was mainly driven by Other International Markets and increased success of our expanded international sales and services infrastructure, led by Japan, China, and Brazil.

Operating Income: Gross profit increased by \$ 50.3 million, representing an improvement in gross profit margins to 54.8% (*prior year period: 53.7%*). We continue to invest in the expansion of our sales and service infrastructure in key markets and in research and development for new products and services. Overall, SG&A expenses increased \$ 18.1 million, while R&D expenses increased \$ 4.4 million. Operating income grew \$ 25.3 million, or 11.9%, compared to the same period last year.

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Net Income: Net income attributable to Sirona shareholders was \$ 175.7 million, an increase of \$ 28.9 million, or 19.7%, over the prior year. The effective tax rate for the fiscal year ended September 30, 2014, was 23.0% (*prior year period: 24.8%*).

Cash Position: At September 30, 2014, the Company had cash and cash equivalents of \$382.8 million and total debt of \$79.5 million, resulting in net cash of \$303.3 million, or an increase of \$137.0 million compared to September 30, 2013.

Significant Factors Affecting Our Operating Performance

Foreign Currency Fluctuations

Although the U.S. Dollar is Sirona's reporting currency, it conducts its business in many currencies, and its functional currencies vary depending on the country of operation. Fluctuations in exchange rates impact Sirona's financial results. The single largest influencing factor is the U.S. Dollar/Euro exchange rate.

Although Sirona does not apply hedge accounting for foreign currency derivatives, it has entered into foreign exchange forward contracts to help mitigate foreign currency exposure. As these agreements are short-term (generally not exceeding six months) and do not cover all underlying exposures, continued fluctuation in exchange rates could materially affect Sirona's results of operations.

Loans made to Sirona under the Senior Facilities Agreement entered into on November 14, 2011, are denominated in the functional currency of the respective borrowers. See "Liquidity and Capital Resources" for a discussion of our Senior Facilities Agreement. However, intra-group loans and other intra-group monetary assets and liabilities are denominated in the functional currency of only one of the parties to the agreements. Where intra-group loans are of a long-term investment nature, the potential fluctuations in exchange rates are reflected within other comprehensive income, whereas exchange rate fluctuations for short-term intra-group loans and other short-term intra-group transactions are recorded in the consolidated statements of income. These fluctuations may be significant in any

period due to changes in exchange rates, especially between the Euro and the U.S. Dollar.

The MDP Transaction and the Exchange

On June 30, 2005, Sirona Holdings Luxco S.C.A. ('Luxco'), a Luxembourg-based holding entity owned by funds managed by Madison Dearborn Partners, Beecken Petty O'Keefe, management and employees of Sirona, obtained control over the Sirona business. The transaction was effected by using new legal entities, Sirona Holding GmbH and its wholly-owned subsidiary Sirona Dental Services GmbH, to acquire 100% of the interest in Sirona Dental Systems Beteiligungs- und Verwaltungs GmbH, the former parent of the Sirona business through a leveraged buy-out transaction (the 'MDP Transaction').

The assets and liabilities acquired in the MDP Transaction and the Exchange were partially stepped up to fair value, and a related deferred tax liability was recorded. The excess of the total purchase price over the fair value of the net assets acquired, including IPR&D, which were expensed at the date of closing of the MDP Transaction and the Exchange, was allocated to goodwill and is subject to periodic impairment testing.

Sirona's cost of goods sold, R&D, SG&A expense, and operating results have been and will continue to be materially affected by depreciation and amortization costs resulting from the step-up to fair value of Sirona's assets and liabilities.

Fluctuations in Operating Results

Sirona's operating results have varied in the past and are likely to vary in the future. These variations result from a number of factors, many of which are substantially outside its control, including:

- the timing of new product introductions by us and our competitors;
- timing of industry tradeshows, particularly the International Dental Show ("IDS");
- changes in relationships with distributors;
- the timing of operational decisions by distributors and end users;
- developments in government reimbursement policies;
- changes in product mix;

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our ability to supply products to meet customer demand;
fluctuations in manufacturing costs;
tax incentives;
currency fluctuations; and
general economic conditions, as well as those specific to the healthcare industry and related industries.

Due to the variations which Sirona has experienced in its operating results, it does not believe that period-to-period comparisons of results of operations of Sirona are necessarily meaningful or reliable as indicators of future performance.

Effective Tax Rate

Sirona's effective tax rate may vary significantly from period to period and, as a global enterprise, can be influenced by many factors. These factors include, but are not limited to, changes in the mix of earnings in countries with differing statutory tax rates (including the result of business acquisitions and dispositions), changes in the valuation of deferred tax assets and liabilities, the results of audits and examinations of previously filed tax returns, tax planning initiatives, tax characteristics of income, changes in exchange rates, as well as the timing and deductibility of expenses for tax purposes. The Company's effective tax rate differs from the U.S. federal statutory rate of 35% primarily as a result of lower effective tax rates on certain earnings outside of the United States.

The company makes no provision for deferred U.S. income taxes on undistributed foreign earnings because as of September 30, 2014, it remained management's intention to continue to indefinitely reinvest such earnings in foreign operations. The distribution of lower-taxed foreign earnings to the U.S. would generally increase the Company's effective tax rate.

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Operations Review	Year ended September 30, 2014		2013		Change		
(\$ in millions)							
REVENUE	\$1,171.1	100.0%	\$1,101.5	100.0%	\$69.6	6.3	%
Dental CAD/CAM Systems	425.8	36.4 %	409.3	37.2 %	16.5	4.0	%
Imaging Systems	399.8	34.1 %	378.0	34.3 %	21.8	5.8	%
Treatment Centers	225.3	19.2 %	210.7	19.1 %	14.6	6.9	%
Instruments	120.2	10.3 %	103.5	9.4 %	16.7	16.1	%
COST OF GOODS SOLD	(529.4)	(45.2 %)	(510.1)	(46.3 %)	(19.3)	3.8	%
GROSS PROFIT ⁽¹⁾	641.7	54.8 %	591.4	53.7 %	50.3	8.5	%
Dental CAD/CAM Systems	291.1	68.4 %	276.7	67.6 %	14.4	5.2	%
Imaging Systems	232.6	58.2 %	221.9	58.7 %	10.7	4.8	%
Treatment Centers	91.5	40.6 %	81.4	38.6 %	10.1	12.4	%
Instruments	51.0	42.4 %	43.2	41.7 %	7.8	18.1	%
Corporate (unallocated)	(24.5)		(31.8)		7.3	(23.0)	%
Selling, general and administrative expense	(350.9)	(30.0 %)	(332.8)	(30.2 %)	(18.1)	5.4	%
Research and development	(64.6)	(5.5 %)	(60.2)	(5.5 %)	(4.4)	7.3	%
Net other operating income (expense)	11.9	1.0 %	14.4	1.3 %	(2.5)	(17.4)	%
OPERATING INCOME	238.1	20.3 %	212.8	19.3 %	25.3	11.9	%
Gain (loss) on foreign currency transactions, net	(0.3)	(0.0 %)	(12.4)	(1.1 %)	12.1	(97.6)	%
Gain (loss) on derivative instruments	(2.5)	(0.2 %)	0.4	0.0 %	(2.9)	(725.0)	%
Interest expense, net	(2.9)	(0.2 %)	(3.4)	(0.3 %)	0.5	(14.7)	%
Other income (expense)	(2.0)	(0.2 %)	0.2	0.0 %	(2.2)	(1,100.0)	%
INCOME BEFORE TAXES	230.4	19.7 %	197.6	17.9 %	32.8	16.6	%
Income tax provision	(53.0)	(4.5 %)	(49.0)	(4.4 %)	(4.0)	8.2	%
NET INCOME	177.4	15.1 %	148.6	13.5 %	28.8	19.4	%

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Less: Net income attributable to noncontrolling interests	(1.7)	(0.1 %)	(1.8)	(0.2 %)	0.1	(5.6 %)
NET INCOME ATTRIBUTABLE TO SIRONA DENTAL SYSTEMS, INC.	\$175.7	15.0 %	\$146.8	13.3 %	\$28.9	19.7 %

(1)
 Percentages refer to the percent of *total* revenues except for segment gross profit information, where percentages refer to segment gross profit margin.

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Revenue

For the fiscal year ended September 30, 2014, revenue was \$ 1,171.1 million, an increase of \$ 69.6 million, or 6.3% (*increased 3.9% Constant Currency*). The strengthening of the Euro throughout most of the fiscal year had a favorable impact on our reported revenue growth across all segments; whereas the weakening of other major currencies (such as Australian Dollar, Brazilian Real, Japanese Yen, etc.) had a negative impact on revenue growth for which no adjustment is made in our Constant Currency measure. Revenue developed by segment and geographically as follows:

By *segment*:

CAD/CAM Systems (*increased 4.0% - up 1.9% Constant Currency*): Revenue growth was driven by international markets, benefiting from new-user demand for our best-in-class products. CAD/CAM generally faced a difficult year-over-year comparison, with prior-year Constant Currency growth of 21.6%. The U.S. also faced a difficult year-over-year comparison due to the successful Omnicam trade-up program in fiscal 2013.

Imaging Systems (*increased 5.8% - up 4.0% Constant Currency*): Revenue growth was driven by increasing demand for our intraoral and Orthophos product lines, particularly in the U.S.

Treatment Centers (*increased 6.9% - up 3.4% Constant Currency*): Treatment Center revenues were boosted by the launch of our new Intego unit in the fourth quarter; however, segment revenues faced an overall challenging year-over-year comparison due to strong revenues in the prior year in connection with the IDS in Cologne, Germany, as well as the last-edition program for our well-renowned M1+ unit.

Instruments (*increased 16.1% - up 12.3% Constant Currency*): Revenue growth was driven by international markets, especially in the hygiene and traditional business. Our increased manufacturing capacity contributed to market expansion.

Geographically:

U.S. (*increased 4.7%*): U.S. revenues continued to benefit from strong, new-user demand for both our CAD/CAM and Imaging products, but faced a difficult overall year-over-year comparison, with prior-year growth of 18.2% benefiting from trade-up programs.

International Markets (*increased 7.1% - increased 3.6% Constant Currency*): Germany, our second-largest individual market, faced a challenging year-over-year comparison due to very strong sales in connection with the IDS in March 2013, with prior-year growth of 23.4% Constant Currency. Strong local sales growth in other international markets (led by Japan, China, and Brazil) was largely offset by the weakening of major currencies in these markets.

Cost of Goods Sold and Gross Profit

Cost of Goods Sold

For the fiscal year ended September 30, 2014, cost of goods sold was \$ 529.4 million, an increase of \$ 19.3 million, or 3.8%. Cost of goods sold as a percentage of sales decreased to 45.2% (*prior year period: 46.3%*). Improvements in product, regional, and channel mix combined with cost reductions to more than offset the headwinds resulting from a strengthening Euro throughout most of the fiscal year and weakening of other currencies such as the Australian Dollar, Brazilian Real, and Japanese Yen.

Cost of goods sold included amortization and depreciation expense resulting from the step-up to fair values of tangible and intangible assets of \$33.3 million (*prior year period: \$37.8 million*). Excluding these amounts, cost of goods sold as a percentage of revenue was 42.4% (*prior year period: 42.9%*).

Gross Profit

Sirona uses gross profit, excluding the impacts of the MDP Transaction and the Exchange, to monitor segment performance. For the fiscal year ended September 30, 2014, gross profit was \$ 641.7 million, an increase of \$ 50.3 million, or 8.5%. Excluding the above amounts in cost of goods sold, gross profit as a percentage of revenue was 57.6% (*prior year period: 57.1%*). The increase in the gross profit margin was mainly due to improvements in product, regional, and channel mix combined with cost reductions, more than offsetting the abovementioned headwinds from foreign exchange effects that cut across all segments.

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By segment, gross profit developed in the fiscal year ended September 30, 2014, compared to the fiscal year ended September 30, 2013 as follows:

CAD/CAM Systems: Segment gross profit benefited from increased revenues. The increase in gross profit margin was mainly driven by product and regional mix and a higher share of new-user sales versus trade ups, the favorable impacts of which were largely offset by negative foreign currency effects.

Imaging Systems: Segment gross profit and gross profit margin benefited from an increase in direct sales and a more favorable product mix within the extra-oral and intra-oral product lines. A less favorable regional mix and price concessions for the extra-oral product lines in certain international markets were only partially offset by cost reductions.

Treatment Centers: Segment gross profit and gross profit margin were boosted by productivity gains and distribution efficiencies achieved but were partially offset by less favorable product and regional mix.

Instruments: Segment gross profit benefited from the strong increase in general sales levels, especially in the hygiene category. The increase in gross profit margin was mainly driven by product and regional mix, partly offset by production inefficiencies and negative foreign currency impacts.

Selling, General, and Administrative

For the fiscal year ended September 30, 2014, SG&A expense was \$ 350.9 million, an increase of \$ 18.1 million, or 5.4%.

The increase in SG&A expense was primarily driven by the increase in sales volume, selling costs related to the introduction of new products, continuing investment in the expansion of our sales and service infrastructure in growth markets, and general inflation.

SG&A expense for the fiscal year ended September 30, 2014 included:

\$12.2 million of other non-cash share-based compensation expense,
\$2.7 million for the CFO Transition, and
\$2.4 million of amortization and depreciation resulting from the step-up to fair values of tangible and intangible assets.

SG&A expense for the fiscal year ended September 30, 2013 included:

\$12.8 million of other non-cash share-based compensation expense (\$9.0 million excluding the impact of the agreements with the former Chairman/CEO and EVP/CFO),
\$8.6 million for the transition agreements with the former Chairman and CEO and the former EVP and CFO (which included a \$3.8 million non-cash charge for the modification of share-based awards),
\$2.5 million of amortization and depreciation resulting from the step-up to fair values of tangible and intangible assets, and
\$3.5 million of costs for the biennial IDS.

Excluding the effects of the transition agreements, SG&A expense increased \$28.6 million, or 9.7% in the fiscal year ended September 30, 2013.

Excluding the above amounts, as a percentage of revenue, SG&A expense was 28.5% (*prior year period: 28.1%*).

Research and Development

R&D expense for the fiscal year ended September 30, 2014, was \$ 64.6 million, an increase of \$ 4.4 million, or 7.3%.

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The increase was mostly driven by additional investment in R&D projects and to a lesser extent the strengthening of the Euro throughout most of the fiscal year. As a percentage of revenue, R&D expense was 5.5% (*prior year period: 5.5%*).

Net Other Operating Income (Loss)

Net other operating income (loss) for the fiscal year ended September 30, 2014, compared to the fiscal year ended September 30, 2013 was as follows:

Net Other Operating Income (Loss)	Year ended	
	September 30, 2014	September 30, 2013
(In millions)		
Income resulting from the amortization of the deferred income related to the Patterson exclusivity payment	\$ 10.0	\$ 10.0
Gain (loss) from patent infringement settlement ⁽¹⁾	-	4.4
Other miscellaneous gain (loss) ⁽²⁾	1.9	-
Net other operating income (loss)	\$ 11.9	\$ 14.4

⁽¹⁾ The gain from patent settlement for the year ended September 30, 2013, represents amounts received for past lost profits in an out-of-court

settlement of a patent defense suit in the normal course of business.

(2) The other miscellaneous gain for the year ended September 30, 2014, represents a gain on disposal of certain business assets.

Gain (Loss) on Foreign Currency Transactions and Derivative Instruments

Foreign Currency Transactions

The loss on foreign currency transactions for the fiscal year ended September 30, 2014 amounted to \$ 0.3 million (prior year period: loss of \$ 12.4 million). The components of these results are as follows:

Gain (Loss) on Foreign Currency Transactions	Year ended	
(In millions)	September 30,	2014 2013
Unrealized non-cash foreign exchange gain (loss) from translation adjustment of deferred income related to the Patterson exclusivity payment	\$(2.0)	\$1.9
Unrealized non-cash foreign exchange gain (loss) on short-term intra-group loans	(0.1)	(0.9)
Gain (loss) on other foreign currency transactions ⁽¹⁾	1.8	(13.4)
Gain (Loss) on Foreign Currency Transactions	\$(0.3)	\$(12.4)

(1) For the fiscal year ended September 30, 2014 and 2013, the gain (loss) on other foreign

currency transactions related to the revaluation of short-term assets and liabilities and realized transactions, both of which were primarily impacted by the fluctuations between the Yen/Euro, Euro/U.S. Dollar, and Real/Euro exchange rates.

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Derivative Instruments

For the fiscal year ended September 30, 2014, the loss on derivative instruments was \$ 2.5 million (*prior year period: gain of \$ 0.4 million*). In both periods, the results related to foreign currency hedges.

Income Tax Provision

For the fiscal year ended September 30, 2014, Sirona recorded a profit before income taxes of \$ 230.4 million (*prior year period: \$ 197.6 million*), an income tax provision of \$ 53.0 million (*prior year period: \$ 49.0 million*), and an effective tax rate of 23.0% (*prior year period: 24.8%*). The income tax provision as of September 30, 2013, included the effect from a local trade tax increase at our principal German operations, which was enacted in and effective beginning in the first quarter of fiscal year 2013. This tax rate change resulted in a tax expense of \$2.2 million from a non-cash remeasurement of deferred tax assets and liabilities. Excluding this amount, the effective tax rate in fiscal year 2013 was 23.7%.

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Operations Review	Year ended				Change		
	September 30,						
	2013		2012				
(\$ in millions)							
REVENUE	\$1,101.5	100.0%	\$979.4	100.0%	\$122.1	12.5	%
Dental CAD/CAM Systems	409.3	37.2 %	334.5	34.2 %	74.8	22.4	%
Imaging Systems	378.0	34.3 %	343.5	35.1 %	34.5	10.0	%
Treatment Centers	210.7	19.1 %	197.1	20.1 %	13.6	6.9	%
Instruments	103.5	9.4 %	102.5	10.5 %	1.0	1.0	%
Corporate (unallocated)			1.8	0.2 %	(1.8)	(100.0)	%
COST OF GOODS SOLD	(510.1)	(46.3 %)	(455.4)	(46.5 %)	(54.7)	12.0	%
GROSS PROFIT ⁽¹⁾	591.4	53.7 %	524.0	53.5 %	67.4	12.9	%
Dental CAD/CAM Systems	276.7	67.6 %	233.8	69.9 %	42.9	18.3	%
Imaging Systems	221.9	58.7 %	199.5	58.1 %	22.4	11.2	%
Treatment Centers	81.4	38.6 %	79.6	40.4 %	1.8	2.3	%
Instruments	43.2	41.7 %	47.0	45.9 %	(3.8)	(8.1	%)
Corporate (unallocated)	(31.8)		(36.0)		4.2	(11.7	%)
Selling, general and administrative expense	(332.8)	(30.2 %)	(295.6)	(30.2 %)	(37.2)	12.6	%
Research and development	(60.2)	(5.5 %)	(52.6)	(5.4 %)	(7.6)	14.4	%
Net other operating income (expense)	14.4	1.3 %	10.0	1.0 %	4.4	44.0	%
OPERATING INCOME	212.8	19.3 %	185.8	19.0 %	27.0	14.5	%
Gain (loss) on foreign currency transactions, net	(12.4)	(1.1 %)	(5.9)	(0.6 %)	(6.5)	110.2	%
Gain (loss) on derivative instruments	0.4	0.0 %	2.0	0.2 %	(1.6)	(80.0	%)
Interest expense, net	(3.4)	(0.3 %)	(3.8)	(0.4 %)	0.4	(10.5	%)
Other income (expense)	0.1	0.0 %	0.2	0.0 %	(0.1)	(50.0	%)
INCOME BEFORE TAXES	197.5	17.9 %	178.3	18.2 %	19.2	10.8	%
Income tax provision	(49.0)	(4.4 %)	(42.7)	(4.4 %)	(6.3)	14.8	%
NET INCOME	148.5	13.5 %	135.6	13.8 %	12.9	9.5	%
Less: Net income attributable to noncontrolling interests	(1.8)	(0.2 %)	(1.8)	(0.2 %)	-	0.0	%
	\$146.7	13.3 %	\$133.8	13.7 %	\$12.9	9.6	%

NET INCOME ATTRIBUTABLE TO SIRONA
DENTAL SYSTEMS, INC.

(1) Percentages refer to the percent of *total* revenues except for segment gross profit information, where percentages refer to segment gross profit margin.

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Revenue

Revenue for the fiscal year ended September 30, 2013, was \$1,101.5 million, an increase of \$122.1 million, or 12.5% (*increased 11.7% Constant Currency*). Revenue developed by segment and geographically as follows:

By segment:

CAD/CAM Systems (*increased 22.4% - up 21.6% Constant Currency*): Segment revenue growth benefited from the Omnicam launch, the delivery of trade-up units, and a generally-increasing demand for products in this segment. Growth was broad-based, but was particularly strong in the U.S. and Germany.

Imaging Systems (*increased 10.0% - up 9.5% Constant Currency*): Sales growth was driven by robust demand for our Orthophos product line, particularly in the U.S. and Germany.

Treatment Centers (*increased 6.9% - up 5.8% Constant Currency*): We continued the above-market growth trajectory. Sales were particularly strong in Germany, benefiting from robust demand for our comfort and standard treatment center product lines, including the last-edition program for our well-renowned M1+ unit.

· **Instruments** (*increased 1.0% - flat Constant Currency*): Sales were on prior-year levels Constant Currency.

Geographically:

U.S. (*increased 18.2%*): Revenues were exceptionally strong and benefited from (i) strong demand for our Imaging and CAD/CAM products, (ii) the impact of the implementation of the Medical Device Tax in 2013 and anticipated changes in tax benefits in the first quarter, (iii) the delivery of Omnicam trade-up units, particularly in the third and fourth quarters, and (iv) the expanded agreement with Patterson.

· **International Markets** (*increased 10.1% - up 9.1% Constant Currency*): Sales growth was particularly driven by an exceptionally strong performance in Germany (up 23.4% Constant Currency), where we experienced strong demand for our CAD/CAM and Imaging products, as well as for the last-edition M1+ treatment center unit. Sales growth in other international markets was overall robust and particularly driven by strong performance in China, Canada,

Russia, and Brazil.

Excluding foreign currency fluctuations, revenue growth was mainly volume and product-mix driven. Prices in general remained stable, with the exception of pricing concessions in certain markets within the Imaging product lines.

Cost of Goods Sold and Gross Profit

Cost of Goods Sold

For the fiscal year ended September 30, 2013, cost of goods sold was \$510.1 million, an increase of \$54.7 million, or 12.0%. Cost of goods sold as a percentage of sales increased slightly to 53.7% (*prior year period: 53.5%*).

Cost of goods sold included amortization and depreciation expense resulting from the step-up to fair values of tangible and intangible assets of \$37.8 million (*prior year period: \$43.9 million*). Excluding these amounts, cost of goods sold as a percentage of revenue was 42.9% (*prior year period: 42.0%*).

Gross Profit

Sirona uses gross profit, excluding the impacts of the MDP Transaction and the Exchange, to monitor segment performance. For the fiscal year ended September 30, 2013, gross profit was \$591.4 million, an increase of \$67.4 million, or 12.9%. Excluding the above amounts in cost of goods sold, gross profit as a percentage of revenue was 57.1% (*prior year period: 58.0%*).

By segment, gross profit developed in the fiscal year ended September 30, 2013 compared to the fiscal year ended September 30, 2012, as follows:

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CAD/CAM Systems: Segment gross profit benefited from the strong increase in sales; however, gross profit margin was below the prior year. The decrease in gross profit margin was mainly due driven by product mix, due to an increasing share of Omnicam sales and from the delivery of trade-ups.

Imaging Systems: Segment gross profit as well as gross profit margin mainly benefited from a strong increase in sales and favorable product mix within the extra-oral and intra-oral product lines.

Treatment Centers: The increase in gross profit was driven by increased volume, whereas the gross profit margin decrease was mainly due to product and regional mix.

Instruments: Segment gross profit and gross profit margin were below the prior-year level. The decrease in gross profit margin was driven by the increased ratio of lower-end handpieces, short-term inefficiencies from the ramp-up of expanded manufacturing capacity, as well as lower hygiene product sales.

Selling, General, and Administrative

For the fiscal year ended September 30, 2013, SG&A expense was \$332.8 million, an increase of \$37.2 million, or 12.6%.

The increase in SG&A expense was primarily driven by the general increase in sales volume and continuing investment in the expansion of our sales and service infrastructure to capitalize on opportunities to gain market share and build up our presence in key growth markets.

SG&A expense for the fiscal year ended September 30, 2013, included:

\$12.8 million of other non-cash share-based compensation expense (\$9.0 million excluding the impact of the agreements with the former Chairman/CEO and ECP/CFO),

\$8.6 million for the transition agreements with the former Chairman and CEO and the former EVP and CFO (which included a \$3.8 million non-cash charge for the modification of share-based awards),

\$2.5 million of amortization and depreciation resulting from the step-up to fair values of tangible and intangible assets, and

\$3.5 million of costs for the biennial IDS.

Excluding the effects of the transition agreements, SG&A expense increased \$28.6 million, or 9.7% in the fiscal year ended September 30, 2013.

SG&A expense for the fiscal year ended September 30, 2012, included:

\$8.4 million of other non-cash share-based compensation expense, and \$2.7 million of amortization and depreciation resulting from the step-up to fair values of tangible and intangible assets.

Excluding the above amounts, as a percentage of revenue, SG&A expense was 28.5% (*prior year period: 29.1%*).

Research and Development

R&D expense for the fiscal year ended September 30, 2013, was \$59.6 million, an increase of \$7.0 million, or 13.2%.

The increase was mainly driven by the timing of projects, particularly in preparation for the biennial IDS in March, where we launched a record 25 new products. As a percentage of revenue, R&D expense was 5.4% (*prior year period: 5.4%*).

Net Other Operating Income (Loss)

Net other operating income (loss) for the fiscal year ended September 30, 2013, compared to the fiscal year ended September 30, 2012 was as follows:

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Net Other Operating Income (Loss)	Year ended September 30,	
(In millions)	2013	2012
Income resulting from the amortization of the deferred income related to the Patterson exclusivity payment	\$10.0	\$10.0
Gain (loss) from patent infringement settlement ⁽¹⁾	4.4	-
Net other operating income (loss)	\$14.4	\$10.0

⁽¹⁾ The gain from patent settlement for the fiscal year ended September 30, 2013, represents amounts received for past lost profits in an out-of-court settlement of a patent defense suit in the normal course of business.

Gain (Loss) on Foreign Currency Transactions and Derivative Instruments

Foreign Currency Transactions

The loss on foreign currency transactions for the fiscal year ended September 30, 2013, amounted to \$12.4 million (*prior year period: loss of \$5.9 million*). The components of these results are as follows:

Gain (Loss) on Foreign Currency Transactions	Year ended September 30,	
(In millions)	2013	2012
Unrealized non-cash foreign exchange gain (loss) from translation adjustment of deferred income related to the Patterson exclusivity payment	\$1.9	\$(2.6)
Unrealized non-cash foreign exchange gain (loss) on short-term intra-group loans	(0.9)	(3.4)
Gain (loss) on other foreign currency transactions ⁽¹⁾	(13.4)	0.1
Gain (Loss) on Foreign Currency Transactions	\$(12.4)	\$(5.9)

⁽¹⁾ For the fiscal years ended September 30, 2013 and 2012, the loss on other foreign currency transactions related to the revaluation of short-term assets and liabilities and realized transactions. For the fiscal year 2013, these were primarily impacted by the weakness of the Yen, as well as fluctuations in the Brazilian Real and South African Rand to the U.S. Dollar in the second half of the fiscal year. For the fiscal year 2012, these were primarily impacted by the fluctuations between the Euro/U.S. Dollar.

Derivative Instruments

For the fiscal year ended September 30, 2013, the gain on derivative instruments was \$ 0.4 million (*prior year period: gain of \$ 2.0 million*). In both periods, the results related to foreign currency hedges.

Income Tax Provision

For the fiscal year ended September 30, 2013, Sirona recorded a profit before income taxes of \$197.6 million (*prior year period: \$178.4 million*), an income tax provision of \$49.0 million (*prior year period: \$42.7 million*), and an effective tax rate of 24.8% (*prior year period: 24.0%*). The income tax provision as of September 30, 2013, included the effect from a local trade tax increase at our principal German operations, which was enacted in and effective beginning in the first quarter of fiscal year 2013. This tax rate change resulted in a tax expense of \$2.2 million from a non-cash remeasurement of deferred tax assets and liabilities.

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Excluding this amount, the effective tax rate in fiscal year 2013 was 23.7%.

LIQUIDITY AND CAPITAL RESOURCES

Summary

Historically, Sirona's principal uses of cash, apart from operating requirements (including research and development efforts), have been for interest payments, debt repayment, and acquisitions. Operating capital expenditures typically are approximately equal to operating depreciation (excluding any effects from the increased amortization and depreciation expense resulting from the step-up to fair values of Sirona's and Schick's assets and liabilities required under purchase accounting). These expenditures may temporarily exceed operating depreciation for larger-scale infrastructure and other investment activities that the Company may undertake from time to time. The Company also uses cash for occasional purchases of treasury shares pursuant to stock repurchase programs.

At September 30, 2014, the Company had cash and cash equivalents of \$382.8 million and total debt of \$79.5 million, resulting in net cash of \$303.3 million. We believe our ability to generate cash from operating activities is one of our fundamental strengths. The near-term outlook for our business remains strong, and we expect to continue generating significant cash flows from operations in the future. The Company typically does not raise capital through issuance of stock; instead, we use debt financing to lower our overall cost of capital and increase our return on shareholders' equity. We believe that our operating cash flows, available cash, and available but unused revolving credit facilities, in combination, provide us with the necessary financial flexibility to fund our working capital needs, research and development efforts, and anticipated capital expenditures for the foreseeable future.

We have significant operations outside of the U.S. and earn a significant portion of our consolidated operating income and income before taxes through our foreign subsidiaries. Cash and cash equivalents of \$ 198.4 million held by our foreign subsidiaries generally are not subject to restrictions prohibiting such amounts from being available in the United States. The distribution of lower-taxed foreign earnings to the United States, however, would generally increase our effective tax rate. It is management's intention to continue to indefinitely reinvest such earnings in foreign operations.

Debt Financing

On November 14, 2011, the Company entered into a senior facilities agreement (the "Senior Facilities Agreement") with Sirona Dental Systems, Inc. and all significant subsidiaries of Sirona as original borrowers and original guarantors, and as of November 16, 2011, Sirona fully repaid its obligations under the Prior Senior Facilities Agreement. Initial borrowings under the Senior Facilities Agreement were used to retire the outstanding borrowings under the Company's previous credit facilities. Please see "Capital Resources - Senior Facilities Agreement" within this section and Note 19 to our consolidated financial statements for a complete description of this Senior Facilities Agreement.

The Senior Facilities Agreement contains restrictive covenants that limit Sirona's ability to make loans, to incur additional indebtedness, and to make disposals, subject to agreed exceptions. The Company has agreed to certain financial debt covenants in relation to the financing. The covenants stipulate that the Company must maintain certain ratios in respect of consolidated total net debt to consolidated adjusted EBITDA. If the Company breaches any of the covenants, the loans will become repayable on demand.

The financial covenants require that the Company maintain a debt coverage ratio ("Debt Cover Ratio") of consolidated total net debt to consolidated adjusted EBITDA ("Consolidated Adjusted EBITDA"), determined on the basis of the last twelve months, of no more than 3.00 to 1. The Company is required to determine its compliance with the covenants as of September 30 and March 31. As of September 30, 2014, the most recent period for which this ratio was calculated, the Company was in compliance:

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Covenants Ratios (In millions, except for ratio amounts)	September 30, 2014	March 31, 2014	September 30, 2013
Consolidated Total Net Debt	\$ (303.3)	\$ (176.8)	\$ (163.4)
Consolidated Adjusted EBITDA	\$ 321.6	\$ 298.3	\$ 294.7
Debt Cover Ratio ⁽¹⁾ as set by covenants (less than or equal to)	N/A 3.00	N/A 3.00	N/A 3.00

⁽¹⁾ Not meaningful in the absence of net debt.

Cash Flows

The following table summarizes our cash flows from operating, investing, and financing activities for each of the periods under report:

Cash Flows (In millions, except for percent amounts)	Year ended September 30,			Change	
	2014	2013	2012	2014 vs. 2013	2013 vs. 2012
Net cash provided by (used in):					
OPERATING activities	\$248.4	\$232.0	\$201.5	\$16.4	7.1 %
INVESTING activities	(81.6)	(106.8)	(52.1)	25.2	23.6 %
FINANCING activities	(10.0)	(38.1)	(345.6)	28.1	73.8 %
Increase (decrease) in cash during the period	\$156.8	\$87.1	\$(196.2)	\$69.7	80.0 %
				\$30.5	15.1 %
				(54.7)	(105.0 %)
				307.5	89.0 %
				\$283.3	(144.4 %)

Net Cash Provided by (Used in) Operating Activities

Net cash provided by operating activities represents net cash from operations, returns on investments, and payments for interest and taxation.

Net cash provided by operating activities was \$248.4 million for the fiscal year ended September 30, 2014 (*prior year period: \$232.0 million*), an increase of \$16.4, or 7.1%.

Influencing factors on the current-period change in operating cash flows were:

Adjustments to reconcile net income to net operating cash flows:

o \$9.2 million less favorable adjustments from derivative instrument and foreign currency transactions due to a significant reduction in net losses versus the prior year.

o \$11.1 million less favorable impacts from net deferred taxes. In the prior year, the acquisition of a technology company and the trade tax increase at our primary facility in Bensheim, Germany, both resulted in favorable adjustments.

o In fiscal 2014, the gain on the sale of business assets resulted in an unfavorable impact of \$1.9 million in the other adjustments.

o The remaining changes only had a minor impact on the decrease in adjustments to reconcile net income to net cash provided by operating activities.

Effect of changes in assets and liabilities:

o Accounts receivable decreased significantly over the prior year due in part to a lower concentration of revenues in the latter part of the fourth quarter in the current year compared to the prior year as well as a shift in customer structure, with increased sales in the fourth quarter to customers with shorter payment terms.

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Changes in trade accounts payable and other assets and liabilities were \$23.9 million less favorable compared to the prior year. This development reflects the timing of supplier payments in the current period as well as the favorable impacts in the prior year resulting from the acquisition of a technology company.

Changes in current income tax liabilities resulted in \$2.1 million more unfavorable adjustments in the current year due to the timing of income tax payments.

Net Cash Provided by (Used in) Investing Activities

Net cash used in investing activities represents cash used for capital expenditures in the normal course of operating activities, financial investments, acquisitions, asset disposals, and divestitures.

Net cash used in investing activities was \$81.6 million for the fiscal year ended September 30, 2014 (*prior year period: \$106.8 million*), a decrease of \$25.2 million, or 23.6%.

The primary contributors to the investing cash outflow in the fiscal year ended September 30, 2014 were:

- the acquisition of the main administrative building in Bensheim, Germany for \$26.7 million,
- the sale of certain business assets for \$11.5 million, and
- software development costs for \$19.8 million,
- and investment in additional machinery for the instruments manufacturing facility in Bensheim, Germany.

For the fiscal year ended September 30, 2013, the primary contributors to the net cash used in investing activities were:

the acquisition of a technology company for \$35.0 million,
the completion of the expansion of the new instruments manufacturing facility in Bensheim, Germany for \$16.9 million,

software development costs for \$13.5 million.

For the fiscal year ended September 30, 2012, the primary contributor to net cash used in investing activities primarily was software development costs for \$11.5 million.

Net Cash Provided by (Used in) Financing Activities

Net cash used in financing activities was \$10.0 million for the fiscal year ended September 30, 2014 (*prior year period: \$38.1 million*), a decrease of \$28.1 million, or 73.8%.

Influencing factors on the current-period change in financing cash flows were:

\$28.1 million fewer purchases of treasury shares compared to the prior period, and \$1.1 million lower proceeds and \$0.3 million higher tax effects from shares issued under share-based compensation plans.

For the fiscal year ended September 30, 2014, primary contributors to net cash used in financing activities were:

\$14.8 million for purchases of treasury shares, less
\$6.8 million of proceeds from shares issued under share-based compensation plans.

For the fiscal year ended September 30, 2013, the primary contributors to net cash used in financing activities were:

\$42.9 million for purchases of treasury shares,
\$2.8 million for the purchase of shares from and dividends paid to noncontrolling interest, less
\$7.9 million of proceeds from shares issued under share-based compensation plans.

For the fiscal year ended September 30, 2012, primary contributors to net cash used in financing activities were:

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\$295.5 million for the repayment of our senior term loans net of new loans received, and \$49.3 million for purchases of treasury shares.

Capital Resources

Senior Facilities Agreement

On November 14, 2011, the Company entered into the Senior Facilities Agreement with Sirona Dental Systems, Inc. and all significant subsidiaries of Sirona as original borrowers and original guarantors. As of November 16, 2011, Sirona fully repaid its obligations under the Prior Senior Facilities Agreement. Initial borrowings under the Senior Facilities Agreement were used to retire the outstanding borrowings under the Company's previous credit facilities.

The Senior Facilities Agreement includes:

- (1) a term loan in an aggregate principal amount of \$75 million ("Facility A Term Loan") available to Sirona or Sirona Dental, as borrower;
- (2) a 120 million Euro revolving credit facility ("Revolving Facility B") available to Sirona Dental Systems GmbH and Sirona Dental Services GmbH, as initial borrowers; and
- (3) a \$100 million revolving credit facility ("Revolving Facility C") available to Sirona or Sirona Dental, as initial borrowers.

The Revolving Facility B is available for borrowing in Euro or any other freely available currency agreed to by the facility agent. The facilities are made available on an unsecured basis. Subject to certain limitations, each European guarantor guarantees the performance of each European borrower, except itself, and each U.S. guarantor guarantees the performance of each U.S. borrower, except itself.

Of the amount borrowed under the Facility A Term Loan, 30% is due on November 16, 2015, and the balance is due on November 16, 2016. The loans under the Senior Facilities Agreement bear interest of EURIBOR, for Euro-denominated loans, and LIBOR for the other loans, plus an initial margin of 160, 85 and 110 basis points for the Facility A Term Loan, Revolving Facility B and Revolving Facility C, respectively.

The Senior Facilities Agreement contains a margin ratchet. Pursuant to this provision, which applies from March 31, 2012 onwards, the applicable margin varies depending on the Company's leverage multiple (i.e. the ratio of consolidated total net debt to consolidated adjusted EBITDA as defined in the Senior Facilities Agreement) between 160 basis points and 215 basis points for the Facility A Term Loan, 85 basis points and 140 basis points for the Revolving Facility B, and 110 basis points and 165 basis points for the Revolving Facility C.

The Senior Facilities Agreement contains restrictive covenants that limit Sirona's ability to make loans, to incur additional indebtedness, and to make disposals, subject to agreed-upon exceptions. The Company has agreed to certain financial debt covenants in relation to the financing. The covenants stipulate that the Company must maintain certain ratios in respect of consolidated total net debt to consolidated adjusted EBITDA. If the Company breaches these covenants, the loans will become repayable on demand.

On November 16, 2011, Sirona entered into 5-year payer interest rate swaps to fully hedge its 3-month LIBOR exposure for the Facility A Term Loan. The terms of the swap reflect the term structure of the underlying loan. The effective nominal interest rate is 1.2775% plus the applicable margin. Settlement of the swaps is required on a quarterly basis.

Debt issuance costs of \$2.8 million were incurred in relation to the financing in November 2011 and have been capitalized as deferred charges and are amortized using the effective interest method over the term of the loans.

As of September 30, 2014 and September 30, 2013, the Facility A Term Loan was fully drawn to the amount of \$75.0 million. The Revolving Facilities B and C remained undrawn as of September 30, 2014 and September 30, 2013.

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The following table summarizes contractual obligations and commercial commitments as of September 30, 2014:

Contractual Obligations and Commercial Commitments	Payments due by period				
	Total	< 1 year	1-3 years	3-5 years	> 5 years
(In millions)					
Long-term debt ⁽¹⁾	\$81.2	\$ 3.4	\$ 77.8	\$ -	\$ -
Capital lease obligations	4.1	1.1	-	-	3.0
Operating lease obligations	27.4	8.9	10.6	5.8	2.1
Pension	31.7	3.6	5.6	6.2	16.3
Purchase commitments ⁽²⁾	20.0	-	20.0	-	-
Total	\$164.4	\$ 17.0	\$ 114.0	\$ 12.0	\$ 21.4

⁽¹⁾ includes expected interest payments and agency/commitment fees

⁽²⁾ represents unconditional purchase commitments with remaining terms in excess of one year

OFF-BALANCE SHEET ARRANGEMENTS

Sirona does not have any off-balance sheet financing arrangements other than its derivatives.

SIGNIFICANT ACCOUNTING POLICIES AND ESTIMATES

Introduction

The preparation of financial statements in conformity with U.S. GAAP requires Sirona to make estimates and assumptions that affect amounts reported in its consolidated financial statements and accompanying notes. These estimates and assumptions are evaluated on an ongoing basis based on historical developments, market conditions, industry trends and other information Sirona believes to be reasonable under the circumstances. There can be no assurance that actual results will conform to Sirona's estimates and assumptions, and that reported results of operations will not be materially adversely affected by the need to make accounting adjustments to reflect changes in its estimates and assumptions from time to time. The following is a discussion of significant accounting policies and estimates important for the understanding of certain events and disclosures for the period under report. For a comprehensive listing and discussion of our significant accounting policies and estimates, please refer to Note 2 "Basis of Presentation and Summary of Significant Accounting Policies" in this report.

Significant Policies and Estimates

Revenue Recognition

The Company's main revenue stream results from the delivery of dental equipment. The Company also enters into revenue arrangements that consist of multiple deliverables of its product and service offerings.

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Additionally, certain products, primarily in our CAD/CAM and Imaging segments, may contain embedded software that functions together with the product to deliver the product's essential functionality.

Revenue, net of related discounts and allowances, is recognized when products or equipment have been shipped, when persuasive evidence of the arrangement exists, the price is fixed or determinable, collectability is reasonably assured, title and risk of loss has passed to customers based on the shipping terms, no significant obligations remain, and allowances for discounts, returns, and customer incentives can be reliably estimated. The Company offers discounts to its distributors if certain conditions are met. Discounts and allowances are primarily based on the volume of products purchased or targeted to be purchased by the individual customer or distributor. Discounts are deducted from revenue at the time of sale or when the discount is offered, whichever is later. The Company estimates volume discounts based on the individual customer's historical and estimated future product purchases. Returns of products, excluding warranty related returns, are infrequent and insignificant. Amounts received from customers in advance of product shipment are classified as deferred income until the revenue can be recognized in accordance with the Company's revenue recognition policy.

Services: Service revenue is generally recognized ratably over the contract term as the specified services are performed. Amounts received from customers in advance of rendering of services are classified as deferred income until the revenue can be recognized upon rendering of those services.

Extended Warranties: The Company offers its customers an option to purchase extended warranties on certain products. The Company recognizes revenue on these extended warranty contracts ratably over the life of the contract. The costs associated with these extended warranty contracts are recognized when incurred.

Multiple-Element Arrangements ("MEAs"): Arrangements with customers may include multiple deliverables, including any combination of equipment, services, and extended warranties. The deliverables included in the Company's MEAs are separated into more than one unit of accounting when (i) the delivered equipment has value to the customer on a stand-alone basis, and (ii) delivery of the undelivered service element(s) is probable and substantially in the control of the Company. Arrangement consideration is then allocated to each unit, delivered or undelivered, based on the relative selling price ("RSP") of each unit of accounting based first on vendor-specific objective evidence ("VSOE") if it exists and then based on estimated selling price ("ESP").

VSOE: In most instances, products are sold separately in stand-alone arrangements. Services are also sold separately through renewals of contracts with varying periods. The Company determines VSOE based on its pricing and

discounting practices for the specific product or service when sold separately, considering geographical, customer, and other economic or marketing variables, as well as renewal rates or stand-alone prices for the service element(s).

ESP: The estimated selling price represents the price at which the Company would sell a product or service if it were sold on a stand-alone basis. When VSOE does not exist for all elements, the Company determines ESP for the arrangement element based on sales, cost and margin analysis, as well as other inputs based on its pricing practices. Adjustments for other market and Company-specific factors are made as deemed necessary in determining ESP.

After separating the elements into their specific units of accounting, total arrangement consideration is allocated to each unit of accounting according to the nature of the revenue as described above and application of the RSP method. Total recognized revenue is limited to the amount not contingent upon future transactions.

Pensions and 401(k) Plan

The Company has defined benefit and defined contribution pension plans and an early retirement plan. Sirona recognizes changes in the funded status of its benefit plans, not yet recognized in the income statement, in other comprehensive income until they are amortized as a component of net periodic benefit cost.

Pension expense is recognized on an accrual basis over the employee's approximate service periods. Defined benefit pension costs are determined by using an actuarial method, which provides for the deferral of actuarial gains and losses (in excess of a specified corridor) that result from changes in assumptions or actual experience differing from that assumed. Costs relating to changes in the benefit plan as well as the transition obligation are amortized. Disclosure of the components of periodic pension cost is also required. When purchase accounting is applied, pension liabilities are recognized for the projected benefit obligation in excess of plan assets.

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The key assumption used in the actuarial calculations for the defined benefit pension plans is the selection of the appropriate discount rate. The discount rate has been selected by reference to market interest rates. The discount rate used reflects the rates available on high quality fixed income investment of appropriate duration at the measurement dates of each year. Fluctuations in market interest rates could impact the amount of pension expense recorded for these plans. The discount rate assumed at September 30, 2014 was 2.30% (*prior year period: 3.40%*). Accordingly, this change in discount rate affected the amount of pension obligation recorded at September 30, 2014.

Plan assets consist of insurance policies with a guaranteed minimum return by the insurance company and an excess profit participation feature for a portion of the benefits. Sirona pays the premiums on the insurance policies but does not manage the investment of the funds; the insurance company makes all decisions on investment of funds, including the allocation to asset groups. The fair value of the plan assets such as equity securities, fixed-income investments, and others is based on the cash surrender values reported by the insurance company.

Contributions made to the defined contribution pension plans and the 401(k) savings plan for U.S. employees are accrued based on the contributions required by the plan.

The Company also has an early retirement plan, Altersteilzeit ('ATZ'), which allows certain German employees who have been accepted into the plan to retire at 60 rather than at the legal retirement age of 67. Eligible employees are those who have attained the age of 59, have completed 12 years of service, and have been accepted to participate in the ATZ plan. Accepted employees join for a period of 2-4 years, during which they work in full active service for 50% of the agreed ATZ plan period, the remaining 50% of the plan period being the passive phase during which the employee does not work. Alternatively, the employee may work for 50% of the time for the entire agreed ATZ plan period. The alternative actually executed is decided via mutual agreement between Sirona and the employee. During the active service period, the employees receive 50% of their salary plus a bonus payment equal to 35% of their salary, and the remaining 50% of their salary, plus a bonus payment equal to 35% of their salary, is paid during the inactive service period. The Company recognizes the salary component of the ATZ plan over the period from the beginning of the ATZ period to the end of the active service period.

Income Taxes

Sirona recognizes deferred tax assets and liabilities based on the differences between the financial statement carrying amounts and the tax basis of assets and liabilities. Sirona regularly reviews its deferred tax assets for recoverability and establishes a valuation allowance, as necessary, based on historical taxable income, projected future taxable income, the expected timing of the reversals of existing temporary differences and the implementation of tax-planning strategies. If Sirona is unable to generate sufficient future taxable income in certain tax jurisdictions, or if there is a material change in the actual effective tax rates or time period within which the underlying temporary differences become taxable or deductible, it could be required to increase its valuation allowance against its deferred tax assets resulting in an increase in its effective tax rate and an adverse impact on operating results. As of September 30, 2014, Sirona had recorded valuation allowances against its deferred tax assets in the amount of \$ 3.1 million (*prior year period: \$ 3.2 million*). Further information on income taxes is provided in Note 8 to the consolidated financial statements appearing elsewhere in this report.

Management believes it is more likely than not that forecasted income, including income that may be generated as a result of certain tax planning strategies, together with the tax effects of the deferred tax liabilities, will be sufficient to fully recover the remaining deferred tax assets. In the event that the Company determines all or part of the net deferred tax assets are not realizable in the future, the Company will make an adjustment to the valuation allowance that would be charged to earnings in the period such determination is made. In addition, the calculation of tax liabilities involves significant judgment in estimating the impact of uncertainties in the application of ASC 740 and other complex tax laws. Resolution of these uncertainties in a manner inconsistent with management's expectations could have a material impact on the Company's financial condition and operating results.

Impairment of Long-Lived and Finite-Lived Assets

Sirona assesses all its long-lived assets for impairment whenever events or circumstances indicate their carrying value may not be recoverable. Sirona's management assesses whether there has been an impairment by comparing anticipated undiscounted future cash flows from operating activities with the carrying value of the asset. The factors considered by Sirona's management in this assessment include operating results, trends and prospects, as well as the effects of obsolescence, demand, competition and other economic factors.

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If an impairment is deemed to exist, management records an impairment charge equal to the excess of the carrying value over the fair value of the impaired assets. This could result in a material charge to earnings.

Impairment of Indefinite-Lived Assets

Goodwill is allocated to each of our reporting units, which we regard to be our operating segments (Dental CAD/CAM Systems, Imaging Systems, Treatment Centers, and Instruments). Sirona assesses goodwill for impairment annually on September 30 unless an event occurs or circumstances change that would more likely than not reduce the fair value of a reporting unit below its carrying value at an earlier date. This evaluation begins with a qualitative assessment to determine if the fair value of its reporting units is more likely than not less than their carrying values. The Company evaluates such qualitative factors as

- (i) the results of the last quantitative impairment assessment,
- (ii) macro- and industry economic conditions such as significant changes in the business and legal climate and competition, and
- (iii) Company-specific assumptions including historical data and experience, operating performance indicators, projections of revenues and expenses and related cash flows, expected long-term growth rates, sale or disposition of a significant portion of the business, the development of its stock price, and other factors.

If we determine that the fair value is more likely than not less than the carrying value, or we decide to bypass the qualitative assessment for a reporting unit, goodwill is tested for impairment under the two-step valuation test. The first step is to estimate the fair value of each reporting unit and compare this estimated fair value with each reporting unit's carrying value. If the fair value is less than the carrying value, additional steps, including an allocation of the estimated fair value to the assets and liabilities of the reporting unit, would be necessary to determine the amount, if any, of goodwill impairment. In this second step, a fair value exercise similar to a business combination would be performed where the individual identifiable assets and liabilities of the reporting unit are valued at fair value with the difference between the fair value of the reporting unit being the implied fair value of goodwill. As of September 30, 2014, based on the qualitative assessment, the Company determined that step one of the impairment test is not required. If we would determine the fair value of a reporting unit, we would use a discounted future cash flow model

to estimate reporting unit fair value. Significant assumptions in a discounted cash flow model would include discount rate, revenue and gross profit margin growth and terminal growth rates based on our judgments, estimates and assumptions.

Sirona evaluates trademarks and in-process research and development (“IPR&D”), which are considered indefinite-lived intangible assets until the associated projects are completed, for impairment at least annually or whenever events or circumstances indicate their carrying value might be impaired. In performing this assessment, Sirona’s management employs a systematic methodology that considers qualitative and quantitative evidence in evaluating whether an impairment is likely to have occurred. These factors include operating results, trends and prospects, as well as the effects of obsolescence, demand, competition and other economic factors. If an impairment is likely to have occurred, an estimate of the fair value of the indefinite-lived intangible assets is performed. The carrying value is considered impaired when it exceeds the fair market value. In such an event, an impairment loss is recognized equal to the amount of that excess. Key assumptions in determining fair value include using the expected discounted cash flows. Once an impairment is determined, an impairment charge is recorded in the consolidated statement of income.

ACCOUNTING STANDARDS ISSUED BUT NOT YET ADOPTED

Please see Note 3 to the consolidated financial statements for any discussions of recently issued accounting standards that have not yet been adopted.

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ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

Sirona is exposed to economic risk from foreign currency exchange rates as well as interest rates. Portions of these risks are hedged; however, they may impact our operating performance.

The following discussion should be read in conjunction with Notes 2, 16, and 19 to Sirona's audited consolidated financial statements appearing elsewhere in this report, which provide further information on Sirona's derivative instruments.

Foreign Currency

Sirona's primary market risk exposure is foreign currency risk. Certain transactions, assets, and liabilities are exposed to foreign currency risk, which can adversely affect our revenues and operating profits. To help mitigate this risk and maximize the economic effectiveness of our foreign currency positions, Sirona enters into forward exchange contracts where practicable.

The Euro is the functional currency for the majority of Sirona's subsidiaries, including its German operations, which are the primary sales and manufacturing operations of Sirona. Sales from other Sirona operations are denominated in various foreign currencies. As a percent of total revenues, sales and operating expenses in Euro, U.S. Dollar, and all other currencies (most importantly: Japanese Yen, Australian Dollar, Chinese Yuan Renminbi, and Brazilian Real) for fiscal year 2014 were approximately as follows:

The most significant influencing factor is the U.S. Dollar/Euro exchange rate. During the periods under review, the U.S. Dollar/Euro exchange rate has fluctuated significantly. The following table presents the relevant U.S. Dollar/Euro exchange rate information for the period(s) under report:

Fluctuations in Exchange Rates (USD/EUR)	Year ended September 30,		
	2014	2013	2012
(In U.S. Dollars)			
Exchange rate used to calculate items in Sirona's financial statements:			
Period-end ⁽¹⁾	\$1.2592	\$1.3499	\$1.2922
Average ⁽²⁾	1.3573	1.3121	1.2992
Fluctuations during the period:			
Low ⁽³⁾	\$1.2592	\$1.2710	\$1.2094
High ⁽⁴⁾	1.3956	1.3639	1.4155

⁽¹⁾ Closing rate as of the balance sheet date.

⁽²⁾ Year-to-date average exchange rate for the fiscal year.

⁽³⁾ Lowest daily exchange rate for the fiscal year.

⁽⁴⁾ Highest daily exchange rate for the fiscal year.

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In order to hedge portions of the transactional exposure to fluctuations in exchange rates, based on forecasted and firmly committed cash flows, Sirona enters into forward foreign currency contracts. Currently, the principal currencies hedged against the Euro are the U.S. Dollar, Japanese Yen, and Australian Dollar. These forward foreign currency contracts are intended to reduce short-term effects of changes in exchange rates. Sirona does not apply hedge accounting to these forward foreign currency contracts.

As of September 30, 2014, we changed our disclosure method of estimating and quantifying our foreign currency risks from a tabular presentation of the assets subject to foreign currency risk without an estimate of the possible effects on future earnings to a value-at-risk (“VAR”) model. We made this change because we believe that use of a VAR method provides our shareholders with a better understanding and estimate of the potential impacts on our earnings due to foreign currency risk.

VAR is defined as the expected gain or loss, for a given confidence level, in the fair value of our portfolio due to positive or adverse market movements over a defined time horizon. The VAR model is not intended to represent actual gains or losses in fair value, including determinations of other-than-temporary losses in fair value in accordance with U.S. GAAP, but is used as a risk estimation and management tool. The distribution of the potential changes in total market value of all transactional asset, and liability positions is computed based on the historical volatilities and correlations among foreign currency exchange rates, assuming normal market conditions.

The VAR is calculated as the total gain or loss that will not be exceeded at the 95.0 percentile confidence level. Several risk factors are not captured in the model, including liquidity, operational, and legal risks. The following table sets forth the one-day VAR for substantially all of our positions for the fiscal years ended September 30, 2014 and 2013:

Foreign Currency Risk		Value at Risk	
		Maximum	
(In millions)	Net Exposure ⁽¹⁾	Gain ⁽²⁾	(Loss) ⁽²⁾

September 30, 2014	\$ 215.5	\$ 1.6	\$ (2.0)
September 30, 2013	141.7	1.6	(1.7)

⁽¹⁾ as of the balance sheet date. Represents the net asset (liability) position exposed to foreign currency risk.

⁽²⁾ for the fiscal years ended

Interest Rate

Sirona is also exposed to interest rate risk associated with short and long-term bank loans bearing variable interest rates. To help mitigate this interest rate risk exposure, Sirona enters into interest rate swap agreements.

On November 16, 2011, Sirona entered into interest rate swaps to fully hedge its interest exposure in connection with the Senior Facilities Agreement dated November 14, 2011. See “Management’s Discussion and Analysis of Financial Conditions and Results of Operations - Long-Term Debt” for further details. Since Sirona’s interest exposure is fully hedged and nearly 100% effective as of September 30, 2014, any sensitivity analysis (such as a hypothetical, instantaneous increase of one percentage point in the interest rates applicable to the variable interest rate debt) would not have had a material impact on interest expense for the periods under report.

Please refer to Note 19 for information on interest rates and scheduled maturities of principal by fiscal year for our outstanding variable-rate indebtedness as of September 30, 2014.

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ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA

The response to this item is included as a separate section of this Annual Report on Form 10-K, beginning on page 59.

ITEM 9. CHANGES IN AND DISAGREEMENTS WITH ACCOUNTANTS ON ACCOUNTING AND FINANCIAL DISCLOSURE

None.

ITEM 9A. CONTROLS AND PROCEDURES

Evaluation of Disclosure Controls and Procedures

Our management, with the participation of our chief executive officer (principal executive officer) and chief financial officer (principal financial officer), evaluated the effectiveness of our disclosure controls and procedures (as defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934), as of September 30, 2014. Based upon this evaluation, our chief executive officer and chief financial officer concluded that, as of September 30, 2014, the Company's disclosure controls and procedures are effective. Our disclosure controls and procedures are designed to ensure that information relating to the Company, including our consolidated subsidiaries, that is required to be disclosed in the reports we file or submit under the Securities Exchange Act of 1934 is recorded, processed, summarized and reported within the time periods specified in Commission's rules and forms, and is accumulated and

communicated to our management, including our principal executive officer and our principal financial officer, as appropriate to allow timely decisions regarding required disclosure.

Management's Annual Report on Internal Control over Financial Reporting

Our management is responsible for establishing and maintaining adequate internal control over the Company's financial reporting (as defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act). Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. 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.

Our management assessed the effectiveness of the Company's internal control over financial reporting as of September 30, 2014. In making this assessment, our management used the criteria set forth by the Committee of Sponsoring Organizations of the Treadway Commission (COSO) in Internal Control Integrated Framework (1992). Based on our assessment, management believes that, as of September 30, 2014, our internal control over financial reporting is effective based on those criteria.

The independent registered public accounting firm, which audited the Company's financial statements included in this Form 10-K, has issued an attestation report on the Company's internal control over financial reporting. Please see attestation report on page 61.

Changes in Internal Control over Financial Reporting

No change in our internal control over financial reporting (as defined in rules 13a-15(f) and 15d-15(f) under the Exchange Act) occurred during the quarter ended September 30, 2014, that has materially affected, or is reasonably likely to materially affect, our internal control over financial reporting.

ITEM 9B. OTHER INFORMATION

None.

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PART III

ITEM 10. DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE

The information required by this item not set forth herein is incorporated by reference to the proxy statement for our 2014 Annual Meeting of Stockholders to be filed with the Securities and Exchange Commission no later than 120 days after the end of the fiscal year (September 30, 2014).

ITEM 11. EXECUTIVE COMPENSATION

The information required by this item not set forth herein is incorporated by reference to the proxy statement for our 2014 Annual Meeting of Stockholders to be filed with the Securities and Exchange Commission no later than 120 days after the end of the fiscal year (September 30, 2014).

ITEM 12. SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT AND RELATED STOCKHOLDER MATTERS

The information required by this item not set forth herein is incorporated by reference to the proxy statement for our 2014 Annual Meeting of Stockholders to be filed with the Securities and Exchange Commission no later than 120 days after the end of the fiscal year (September 30, 2014).

ITEM 13. CERTAIN RELATIONSHIPS AND RELATED TRANSACTIONS, AND DIRECTOR INDEPENDENCE

The information required by this item not set forth herein is incorporated by reference to the proxy statement for our 2014 Annual Meeting of Stockholders to be filed with the Securities and Exchange Commission no later than 120 days after the end of the fiscal year (September 30, 2014).

ITEM 14. PRINCIPAL ACCOUNTANT FEES AND SERVICES

The information required by this item not set forth herein is incorporated by reference to the proxy statement for our 2014 Annual Meeting of Stockholders to be filed with the Securities and Exchange Commission no later than 120 days after the end of the fiscal year (September 30, 2014).

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PART IV

ITEM 15. EXHIBITS, FINANCIAL STATEMENT SCHEDULES

(a) (1) Financial Statements, See Financial Information Table of Contents on Page 58.

(b) The following Exhibits are included in this report:

Exhibit No.	Item Title
2.1	Exchange Agreement, by and among Sirona Holdings Luxco S.C.A, Blitz 05-118 GmbH and Schick Technologies, Inc., dated September 25, 2005 (incorporated by reference to Exhibit 99.1 to Form 8-K, filed on September 26, 2005)
2.2	Amendment No. 1 to Exchange Agreement, dated May 11, 2006 (incorporated by reference to Exhibit 99.1 to Form 8-K, filed on May 16, 2006)
3.1	Amended and Restated Certificate of Incorporation of the Company (incorporated by reference to Exhibit 3.1 to the Company's Registration Statement on Form S-1, File No. 333-33731, filed on June 30, 1997)
3.2	Certificate of Amendment to Amended and Restated Certificate of Incorporation of the Company (incorporated by reference to Exhibit 3.2 to Form 8-K filed on June 20, 2006)
3.3	Certificate of Amendment to Amended and Restated Certificate of Incorporation of the Company (incorporated by reference to Exhibit 3.1 to Form 8-K filed on February 24, 2014)
3.4	Bylaws of the Company effective as of September 20, 2010 (incorporated by reference to Exhibit 3.2 to Form 8-K, filed on September 23, 2010)

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- 4.1 Form of Common Stock certificate of the Company (incorporated by reference to Exhibit 4.4 to the Company's Registration Statement on Form S-3, File No. 333-153092, filed on August 20, 2008)
- 10.1 1996 Employee Stock Option Plan, as amended (incorporated by reference to Exhibit 10.1 to Form 10-K, filed on July 13, 2001)†
- 10.2 Amendment to 1996 Employee Stock Option Plan (incorporated by reference to the Company's definitive proxy statement on Schedule 14A, filed on May 16, 2006)†
- 10.3 1997 Stock Option Plan for Non-Employee Directors, as amended (incorporated by reference to Exhibit 10.2 to Form 10-K, filed on June 18, 2003)†
- 10.4 Sirona Dental Systems, Inc. Equity Incentive Plan (incorporated by reference to the Company's definitive proxy statement on Schedule 14A, filed on January 26, 2007)†
- 10.5 Form of Stock Option Notice under Sirona Dental Systems, Inc. Equity Incentive Plan (incorporated by reference Exhibit 10.2 to Form 8-K filed on February 28, 2007)†
- 10.6 Distributorship Agreement, dated April 6, 2000, by and between Schick Technologies, Inc. and Patterson Dental Company (incorporated by reference to Exhibit 10.34 to Form 10-K, filed on June 29, 2000)**
- 10.7 Amendment No. 1 to Distributorship Agreement, dated July 1, 2005 by and between Schick Technologies, Inc. and Patterson Dental Company (incorporated by reference to Exhibit 10.1 to Form 10-Q/A, filed on March 24, 2006)**
- 10.8 Consulting and Non-Competition Agreement between Schick Technologies, Inc. and David B. Schick, dated May 7, 2004 (incorporated by reference to Exhibit 10.33 to Form 10-K, filed on June 25, 2004)
- 10.9 Transaction Services Agreement by and between Blitz F04-506 GmbH, Sirona Dental Services GmbH & Co KG, Sirona Dental Systems GmbH, MDP IV Offshore GP, LP and Harry M. Jansen Kraemer, Jr., dated July 6, 2005 (incorporated by reference to Exhibit 10.7 to Form 10-K, filed on December 11, 2006)
- 10.10 Registration Agreement between the Company and Luxco, dated as of June 20, 2006 (incorporated by reference to Exhibit 2.1 to Form 8-K filed on June 20, 2006)
- 10.11 Employment Agreement between the Company and Jeffrey T. Slovin, dated as of June 14, 2006 (incorporated by reference to Exhibit 2.2 to Form 8-K filed on June 20, 2006)†

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- 10.12 Employment Agreement between the Company and Michael Stone, dated as of June 14, 2006 (incorporated by reference to Exhibit 2.3 to Form 8-K filed on June 20, 2006)†
- 10.13 Transition and Severance Agreement between the Company and Zvi Raskin, dated as of June 14, 2006 (incorporated by reference to Exhibit 2.4 to Form 8-K filed on June 20, 2006)†
- 10.14 Employment Agreement between Sirona Beteiligungs- und Verwaltungsgesellschaft mbH (represented by its shareholder Sirona Dental Systems SARL) and Jost Fischer, dated as of January 25, 2002 (incorporated by reference to Exhibit 10.5 to Form 10-Q, filed on August 9, 2006)†
- 10.15 Employment Agreement between Sirona Beteiligungs- und Verwaltungsgesellschaft mbH (represented by its shareholder Sirona Dental Systems SARL) and Simone Blank, dated as of June 27, 2001 (incorporated by reference to Exhibit 10.6 to Form 10-Q, filed on August 9, 2006)†
- 10.16 Consolidated and Restated Amendment to Distributorship Agreement between Sirona Dental Systems GmbH and Patterson Companies, Inc. (incorporated by reference to Exhibit 10.8 to Form 10-Q, filed on August 9, 2006)**
- 10.17 Senior Facilities Agreement (incorporating amendments made on December 5, 2006 and January 19, 2007) among Sirona Dental Systems, Inc., Schick Technologies, Inc., Sirona Dental Systems GmbH, Sirona Dental Services GmbH, Sirona Dental Systems LLC, Sirona Holding GmbH, Sirona Immobilien GmbH, J.P. Morgan PLC, UBS Limited, JPMorgan Chase Bank, N.A., and J.P. Morgan Europe Limited, dated November 22, 2006 (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 10, 2007)
- 10.18 Description of the Sirona Dental Systems, Inc. EVA Plan (incorporated by reference to Exhibit 10.18 to Form 10-K filed on December 7, 2007)†
- 10.19 Employment Agreement between Schick Technologies, Inc. and Jeffrey T. Slovin, dated June 9, 2004 (superseded by the employment agreement dated June 20, 2006 (the “2006 employment agreement”) incorporated by reference as Exhibit 10.11 to this Form 10-K, except for the bonus information contained in Section IV referenced in the 2006 employment agreement)†
- 10.20 Company’s 2008 Executive Bonus Plan (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 8, 2008)†
- 10.21 Company’s 2009 Executive Bonus Plan (incorporated by reference to Exhibit 10.21 to Form 10-K, filed on December 4, 2008)†
- 10.22 Amended and Restated Service Agreement between Sirona Dental GmbH, the Company and Jost Fischer, dated as of December 2, 2008 (superseding an Executive Service Agreement between Sirona Dental GmbH and Jost Fischer, dated as of October 10, 2007, which superseded the Employment Agreement between Sirona Beteiligungs- und Verwaltungsgesellschaft mbH (represented by its shareholder Sirona Dental Systems SARL) and Jost Fischer, dated as of January 25, 2002) (incorporated by reference to Exhibit 10.22 to Form 10-K, filed on December 4, 2008)†
- 10.23 Amended and Restated Service Agreement between Sirona Dental GmbH, the Company and Simone Blank, dated as of December 2, 2008 (superseding an Executive Service Agreement between Sirona Dental GmbH and

Simone Blank, dated as of October 1, 2007, which superseded the Employment Agreement between Sirona Beteiligungs- und Verwaltungsgesellschaft mbH (represented by its shareholder Sirona Dental Systems SARL) and Simone Blank, dated as of June 27, 2001) (incorporated by reference to Exhibit 10.23 to Form 10-K, filed on December 4, 2008)†

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- Amendment to Employment Agreement, dated as of December 2, 2008, between the Company and Jeffrey T. Slovin (amending the Employment Agreement between the Company and Jeffrey T. Slovin, dated as of June 14, 2006 and superseding the Employment Agreement between the Company and Jeffrey T. Slovin dated as of June 9, 2004) (incorporated by reference to Exhibit 10.24 to Form 10-K, filed on December 4, 2008)†
- 10.24 Sirona Dental Systems, Inc. Equity Incentive Plan, as amended (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on March 3, 2009)†
- 10.25 Schick Technologies, Inc. 1996 Stock Option Plan, as amended (incorporated by reference to Exhibit 10.2 to Form 8-K, filed on March 3, 2009)†
- 10.26 Renewal Letter Agreement, dated as of May 4, 2009, between Sirona Dental Services GmbH, a corporation organized under the laws of Germany (“Sirona GmbH”) and Sirona Holdings Luxco S.C.A., a société en commandite par actions organized under the laws of the Grand Duchy of Luxembourg (“Luxco”), to the Advisory Services Agreement dated October 1, 2005 between Sirona GmbH and Luxco, together with the Assignment and Assumption Agreement dated May 4, 2009 among Sirona GmbH, Sirona Dental Systems, Inc. and Luxco (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 5, 2009)
- 10.27 Form of Restricted Stock Unit Agreement for December 8, 2009 restricted stock unit grants (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on December 11, 2009)†
- 10.28 Amendment to Distributorship Agreement, dated May 5, 2010, by and between Schick Technologies, Inc. and Patterson Companies, Inc. (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 5, 2010)**
- 10.29 Amendment No. 2 to Amended and Restated Employment Agreement, dated as of September 20, 2010, between the Company and Jeffrey T. Slovin (amending the Employment Agreement between the Company and Jeffrey T. Slovin, dated as of June 14, 2006) (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on September 23, 2010)†
- 10.30 Employment Agreement, dated as of September 13, 2007, as amended on October 15, 2008, by and between Sirona Dental GmbH and Walter Petersohn (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on September 23, 2010)†
- 10.31 Supplement Agreement to Service Agreement between Sirona Dental GmbH, the Company and Jost Fischer, dated as of November 15, 2010, as amended by the Amended and Restated Service Agreement between Sirona Dental GmbH, the Company and Jost Fischer, dated as of December 2, 2008 (incorporated by reference to Exhibit 10.32 to Form 10-K, filed on November 18, 2010)†**
- 10.32 Supplement Agreement to Service Agreement between Sirona Dental GmbH, the Company and Simone Blank, dated as of November 15, 2010, as amended by the Amended and Restated Service Agreement between Sirona Dental GmbH, the Company and Simone Blank, dated as of December 2, 2008 (incorporated by reference to Exhibit 10.33 to Form 10-K, filed on November 18, 2010)†**
- 10.33 Amendment to Consolidated and Restated Amendment to Distributorship Agreement, dated May 3, 2011, between Patterson Companies, Inc. and Sirona Dental Systems GMBH (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 6, 2011)**
- 10.34

Term and Revolving Facilities Agreement between, among others, Sirona Dental Systems, Inc., Schick Technologies, Inc., Sirona Dental Systems, LLC, Sirona Dental Services GmbH, Sirona Dental Systems GmbH, 10.35 Sirona Immobilien GmbH, Sirona Technologie GmbH & Co. KG, JPMorgan Limited, UniCredit Bank AG and J.P. Morgan Europe Limited, dated November 14, 2011. (incorporated by reference to Exhibit 10.35 to Form 8-K, filed on November 18, 2011)

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	Senior Facilities Agreement, dated November 4, 2011, by and among Sirona Dental Systems, Inc., J.P.
10.36	Morgan Limited, Unicredit Bank AG and J.P. Morgan Europe Limited. (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on November 18, 2011)
10.37	Amended and Restated U.S. Distributorship Agreement, dated May 31, 2012, by and between Patterson Companies, Inc. and Sirona Dental Systems, Inc. (incorporated by reference to Exhibit 10.1 to Form 8-K/A, filed on July 12, 2012)
10.38	Amended and Restated U.S. CAD-CAM Distributorship Agreement, dated May 31, 2012, by and between Patterson Companies, Inc. and Sirona Dental Systems GmbH. (incorporated by reference to Exhibit 10.2 to Form 8-K/A, filed on July 12, 2012)
10.39	Letter Amendment to Amended and Restated Employment Agreement, dated as of October 1, 2012, between Sirona Dental Systems, Inc. and Jeffrey T. Slovin (incorporated by reference to Exhibit 10.1 to Form 8-K, filed on October 5, 2012)†
10.40	Transition Agreement by and between Sirona Dental GmbH, Sirona Dental Systems, Inc. and Jost Fischer, dated November 16, 2012 (incorporated by reference to Exhibit 10.39 to Form 10-K, filed on November 16, 2012)†
10.41	Employment Contract between Sirona Dental Services GmbH and Rainer Berthan, dated February 20, 2012 (incorporated by reference to Exhibit 10.40 to Form 10-K, filed on November 16, 2012)†
10.42	Amendment to Amended and Restated Employment Agreement, dated as of May 7, 2013, between Sirona Dental Systems, Inc. and Jeffrey T. Slovin (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on May 10, 2013)†
10.43	Executive Employment Agreement, dated July 29, 2013, by and between Sirona Dental Systems, Inc. and Ulrich Michel (incorporated by reference to Exhibit 10.2 to Form 10-Q, filed on August 2, 2013)†
10.44	Separation Agreement, dated August 1, 2013, by and between Sirona Dental GmbH, Sirona Dental Systems, Inc. and Simone Blank (incorporated by reference to Exhibit 10.1 to Form 10-Q, filed on August 2, 2013)†
14.1	Code of Ethics (incorporated by reference to Exhibit 14.1 to the Company's Annual Report on Form 10-K, filed on June 25, 2004)
16.1	Letter from Grant Thornton LLP to the Securities and Exchange Commission confirming statements made about it by Company in connection with changes to the Company's certifying accountant (incorporated by reference to Exhibit 16.1 to Form 8-K, filed on June 26, 2006)
21.1	List of Subsidiaries of Company*
23.1	Consent of Independent Registered Public Accounting Firm*
31.1	Certification of Principal Executive Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002*
31.2	Certification of Principal Financial Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002*
32.1	Section 1350 Certification of Chief Executive Officer*
32.2	Section 1350 Certification of Chief Financial Officer*
101.INS	XBRL Instance Document***

101.SCHXBRL Taxonomy Extension Schema Document***

101.CALXBRL Taxonomy Extension Calculation Linkbase Document***

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101.DEF XBRL Taxonomy Extension Definition Linkbase Document***

101.LAB XBRL Taxonomy Labels Linkbase Document***

101.PRE XBRL Taxonomy Extension Presentation Linkbase Document***

† Compensatory plan or arrangement

* Filed herewith

** Certain information in this exhibit has been omitted and filed separately with the Securities and Exchange Commission pursuant to a confidential treatment request under Rule 24b-2 promulgated under the Securities Exchange Act of 1934, as amended.

*** Attached as Exhibit 101 to this report are the following documents formatted in XBRL (eXtensible Business Reporting Language): (i) Consolidated Statements of Income for the years ended September 30, 2014, 2013 and 2012, (ii) Consolidated Statements of Comprehensive Income for the years ended September 30, 2014, 2013 and 2012, (iii) Consolidated Balance Sheets as of September 30, 2014 and 2013, (iv) Consolidated Statements of Shareholders' Equity for the years ended September 30, 2014, 2013 and 2012, (v) Consolidated Statements of Cash Flows for the years ended September 30, 2014, 2013 and 2012, and (vi) Notes to Consolidated Condensed Financial Statements.

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SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this Annual Report on Form 10-K to be signed on its behalf by the undersigned, thereunto duly authorized.

Dated: November 21, 2014 SIRONA DENTAL SYSTEMS, INC.

By: /s/ JEFFREY T. SLOVIN

Jeffrey T. Slovin

President, Chief Executive Officer, and Director

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

SIGNATURE	TITLE	DATE
/s/ JEFFREY T. SLOVIN Jeffrey T. Slovin	President, Chief Executive Officer, and Director (Principal Executive Officer)	November 21, 2014
/s/ ULRICH MICHEL Ulrich Michel	Executive Vice President and Chief Financial Officer (Principal Financial Officer)	November 21, 2014
/s/ STEPHAN MITSDOERFFER Stephan Mitsdoerffer	Chief Accounting Officer (Principal Accounting Officer)	November 21, 2014

/s/ THOMAS JETTER Thomas Jetter	Chairman of the Board and Director	November 21, 2014
/s/ DAVID BEECKEN David Beecken	Director	November 21, 2014
/s/ WILLIAM K. HOOD William K. Hood	Director	November 21, 2014
/s/ ARTHUR D. KOWALOFF Arthur D. Kowaloff	Director	November 21, 2014
/s/ HARRY M. JANSEN KRAEMER, JR. Harry M. Jansen Kraemer, Jr.	Director	November 21, 2014
/s/ TIMOTHY P. SULLIVAN Timothy P. Sullivan	Director	November 21, 2014

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REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

The Board of Directors

Sirona Dental Systems, Inc.:

We have audited the accompanying consolidated balance sheets of Sirona Dental Systems, Inc. and subsidiaries as of September 30, 2014 and 2013, and the related consolidated statements of income, comprehensive income, shareholders' equity, and cash flows for each of the years in the three-year period ended September 30, 2014. These consolidated financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these consolidated 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 consolidated financial statements referred to above present fairly, in all material respects, the financial position of Sirona Dental Systems, Inc. and subsidiaries as of September 30, 2014 and 2013, and the results of their operations and their cash flows for each of the years in the three-year period ended September 30, 2014 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), Sirona Dental Systems, Inc.'s internal control over financial reporting as of September 30, 2014, based on criteria established in Internal Control—Integrated Framework (1992) issued by the Committee of Sponsoring Organizations of the Treadway Commission (COSO), and our report dated November 21, 2014 expressed an unqualified opinion on the effectiveness of Sirona Dental Systems, Inc.'s internal control over financial reporting.

/s/ KPMG AG Wirtschaftsprüfungsgesellschaft

Frankfurt, Germany

November 21, 2014

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REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

The Board of Directors
Sirona Dental Systems, Inc.:

We have audited Sirona Dental Systems, Inc.'s internal control over financial reporting as of September 30, 2014, based on criteria established in Internal Control—Integrated Framework (1992) issued by the Committee of Sponsoring Organizations of the Treadway Commission (COSO). Sirona Dental Systems, Inc.'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 under Item 9A. 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, and testing and evaluating the design and operating effectiveness of internal control based on the assessed risk. Our audit also included 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, Sirona Dental Systems, Inc. maintained, in all material respects, effective internal control over financial reporting as of September 30, 2014, based on criteria established in Internal Control—Integrated Framework (1992) issued by COSO.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States), the consolidated balance sheets of Sirona Dental Systems, Inc. and subsidiaries as of September 30, 2014 and 2013, and the related consolidated statements of income, comprehensive income, shareholders' equity, and cash flows for each of the years in the three-year period ended September 30, 2014, and our report dated November 21, 2014 expressed an unqualified opinion on those consolidated financial statements.

/s/ KPMG AG Wirtschaftsprüfungsgesellschaft

Frankfurt, Germany
November 21, 2014

Table of Contents**SIRONA DENTAL SYSTEMS, INC.****AND SUBSIDIARIES****FORM 10-K****FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2014****CONSOLIDATED STATEMENTS OF INCOME**

(In millions, except for share and per share amounts)	Notes	Year ended September 30,		
		2014	2013	2012
REVENUE		\$ 1,171.1	\$ 1,101.5	\$ 979.4
Cost of goods sold		(529.4)	(510.1)	(455.4)
GROSS PROFIT		641.7	591.4	524.0
Selling, general and administrative expense		(350.9)	(332.8)	(295.6)
Research and development expense		(64.6)	(60.2)	(52.6)
Net other operating income (loss)	5	11.9	14.4	10.0
OPERATING INCOME		238.1	212.8	185.8
Gain (loss) on foreign currency transactions		(0.3)	(12.4)	(5.9)
Gain (loss) on derivative instruments	6	(2.5)	0.4	2.0
Interest income (expense)		(2.9)	(3.4)	(3.8)
Other income (expense)		(2.0)	0.1	0.2
INCOME BEFORE TAXES		230.4	197.5	178.3
Income tax benefit (expense)	8	(53.0)	(49.0)	(42.7)
NET INCOME		177.4	148.5	135.6
Net (income) loss attributable to noncontrolling interests		(1.7)	(1.8)	(1.8)
NET INCOME ATTRIBUTABLE TO SIRONA DENTAL SYSTEMS, INC.		\$ 175.7	\$ 146.7	\$ 133.8
INCOME PER SHARE				
<i>(attributable to Sirona Dental Systems, Inc. common shareholders)</i>	9			
Basic		\$ 3.18	\$ 2.67	\$ 2.41

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Diluted	\$3.13	\$2.61	\$2.36
Weighted average shares - basic	55,269,606	54,979,044	55,524,188
Weighted average shares - diluted	56,203,970	56,213,992	56,755,396

The accompanying Notes are an integral part of these financial statements.

Table of Contents**SIRONA DENTAL SYSTEMS, INC.****AND SUBSIDIARIES****FORM 10-K****FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2014****CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME**

(In millions)	Notes	Year ended September 30,		
		2014	2013	2012
NET INCOME		\$177.4	\$148.5	\$135.6
OTHER COMPREHENSIVE INCOME (LOSS), NET OF TAX	11			
Cumulative translation adjustment		(88.5)	36.7	(50.0)
Derivative financial hedging instruments		0.3	0.6	(1.3)
Unrecognized elements of pension cost		6.8	0.9	10.2
TOTAL OTHER COMPREHENSIVE INCOME (LOSS)		(81.4)	38.2	(41.1)
TOTAL COMPREHENSIVE INCOME (LOSS)		96.0	186.7	94.5
Comprehensive (income) loss attributable to noncontrolling interests		(1.7)	(1.6)	(1.8)
COMPREHENSIVE INCOME (LOSS) ATTRIBUTABLE TO SIRONA DENTAL SYSTEMS, INC. SHAREHOLDERS		\$94.3	\$185.1	\$92.7

The accompanying Notes are an integral part of these financial statements.

Table of Contents**SIRONA DENTAL SYSTEMS, INC.****AND SUBSIDIARIES****FORM 10-K****FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2014****CONSOLIDATED BALANCE SHEETS**

(In millions, except for share and par value amounts)	Notes	September 30, 2014	2013
ASSETS			
CURRENT ASSETS			
Cash and cash equivalents		\$382.8	\$241.7
Restricted cash		0.8	0.7
Accounts receivable, <i>net of allowance for doubtful accounts of \$2.4 and \$1.6, respectively</i>		115.6	145.2
Inventories, net	13	123.4	109.2
Deferred tax assets	8	29.7	31.4
Prepaid expenses and other current assets		26.8	32.6
Income tax receivable	8	8.0	2.3
TOTAL CURRENT ASSETS		687.1	563.1
Property, plant and equipment, <i>net of accumulated depreciation of \$171.3 and \$156.7, respectively</i>		221.0	182.7
Goodwill		629.3	672.1
Intangible assets, <i>net of accumulated amortization of \$511.3 and \$503.1, respectively</i>		252.8	301.7
Other non-current assets		5.3	5.6
Deferred tax assets	8	15.5	13.2
TOTAL ASSETS		\$1,811.0	\$1,738.4
LIABILITIES AND SHAREHOLDERS' EQUITY			
CURRENT LIABILITIES			
Trade accounts payable		\$59.9	\$73.2
Short-term financial liabilities		1.5	0.4
Income taxes payable	8	6.1	9.3

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Deferred tax liabilities	8	2.2	0.7
Accrued liabilities and deferred income		167.6	162.5
TOTAL CURRENT LIABILITIES		237.3	246.1
Long-term financial liabilities		78.0	75.0
Deferred tax liabilities	8	111.8	131.5
Other non-current liabilities		25.1	27.4
Pension related provisions		71.7	66.0
Deferred income		25.9	34.4
TOTAL LIABILITIES		549.8	580.4
COMMITMENTS & CONTINGENCIES			
SHAREHOLDERS' EQUITY			
Preferred stock (<i>\$0.01 par value; 5,000,000 shares authorized; none issued and outstanding</i>)		-	-
Common stock (<i>\$0.01 par value; 95,000,000 shares authorized; 57,776,336 shares issued; 55,364,617 shares outstanding at Sep. 30, 2014; 57,213,615 shares issued; 54,999,436 shares outstanding at Sept. 30, 2013</i>)		0.6	0.6
Additional paid-in capital		697.9	674.2
Treasury stock, <i>at cost</i> 2,411,719 shares held at cost at Sep. 30, 2014; 2,214,179 shares held at cost at Sept. 30, 2013		(126.8)	(112.0)
Retained earnings		759.9	584.2
Accumulated other comprehensive income (loss)	11	(72.8)	8.6
TOTAL SIRONA DENTAL SYSTEMS, INC. SHAREHOLDERS' EQUITY		1,258.8	1,155.6
NONCONTROLLING INTERESTS		2.4	2.4
TOTAL SHAREHOLDERS' EQUITY		1,261.2	1,158.0
TOTAL LIABILITIES & SHAREHOLDERS' EQUITY		\$1,811.0	\$1,738.4

The accompanying Notes are an integral part of these financial statements.

Table of Contents**SIRONA DENTAL SYSTEMS, INC.****AND SUBSIDIARIES****FORM 10-K****FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2014****CONSOLIDATED STATEMENTS OF SHAREHOLDERS' EQUITY**

(In millions, except for share amounts)	Notes	September 30, 2014	2013	2012
SIRONA DENTAL SYSTEMS, INC. SHAREHOLDERS' EQUITY				
NUMBER OF COMMON SHARES ISSUED AND OUTSTANDING				
Balance at beginning of period		54,999,436	55,051,673	55,815,323
Issuance of common stock upon exercise of options		562,721	615,570	305,625
Purchase of treasury stock		(197,540)	(667,807)	(1,069,275)
Balance at end of period		55,364,617	54,999,436	55,051,673
COMMON SHARE CAPITAL		\$0.6	\$0.6	\$0.6
ADDITIONAL PAID-IN CAPITAL				
Balance at beginning of period		674.2	650.2	636.6
Issuance of common stock upon exercise of options and net effect of vesting of RSUs/PSUs		6.8	7.9	3.9
Stock compensation		12.2	12.8	8.6
Tax effect of stock options exercised and net effect of vesting of RSUs/PSUs		4.7	3.9	0.4
Purchase of shares from noncontrolling interests		-	(0.6)	0.7
Balance at end of period		697.9	674.2	650.2
TREASURY STOCK				

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Balance at beginning of period	(112.0)	(69.1)	(19.8)
Purchase of treasury stock (at cost)	(14.8)	(42.9)	(49.3)
Balance at end of period	(126.8)	(112.0)	(69.1)
RETAINED EARNINGS						
Balance at beginning of period	584.2		437.5		303.7	
Net income (loss)	175.7		146.7		133.8	
Balance at end of period	759.9		584.2		437.5	
ACCUMULATED OTHER COMPREHENSIVE INCOME						
(LOSS)		11				
Balance at beginning of period	8.6		(29.8)	11.3	
Other comprehensive income (loss)	(81.4)	38.4)	(41.1)
Balance at end of period	(72.8)	8.6)	(29.8)
TOTAL SIRONA DENTAL SYSTEMS, INC. SHAREHOLDERS' EQUITY	\$ 1,258.8		\$ 1,155.6		\$ 989.4	
EQUITY ATTRIBUTABLE TO NONCONTROLLING INTERESTS						
Balance at beginning of period	\$2.4		\$3.0		\$3.6	
Purchase of shares from noncontrolling interests	-		(0.8)	(0.7)
Dividend distribution to noncontrolling interests	(1.5)	(1.4)	(1.7)
Net income (loss) attributable to noncontrolling interests	1.7		1.8		1.8	
Cumulative translation adjustment	(0.2)	(0.2)	-	
TOTAL EQUITY ATTRIBUTABLE TO NONCONTROLLING INTERESTS	\$2.4		\$2.4		\$3.0	
TOTAL SHAREHOLDERS' EQUITY	\$ 1,261.2		\$ 1,158.0		\$ 992.4	

The accompanying Notes are an integral part of these financial statements.

Table of Contents**SIRONA DENTAL SYSTEMS, INC.****AND SUBSIDIARIES****FORM 10-K****FOR THE FISCAL YEAR ENDED SEPTEMBER 30, 2014****CONSOLIDATED STATEMENTS OF CASH FLOWS**

	Year ended September 30,		
(In millions)	2014	2013	2012
OPERATING ACTIVITIES			
NET INCOME	\$177.4	\$148.5	\$135.6
ADJUSTMENTS TO RECONCILE NET INCOME TO NET CASH PROVIDED BY (USED IN) OPERATING ACTIVITIES			
Depreciation and amortization	76.3	75.6	77.7
(Gain) loss on derivative instruments and foreign currency transactions	2.8	12.0	3.9
Deferred income taxes	(13.8)	(2.7)	(17.3)
Share-based compensation expense	12.2	12.8	8.6
Other adjustments	(1.3)	0.6	0.6
TOTAL ADJUSTMENTS TO RECONCILE NET INCOME TO OPERATING CASH FLOWS	76.2	98.3	73.5
CHANGES IN ASSETS AND LIABILITIES			
Accounts receivable	20.5	(13.2)	(38.3)
Inventories	(24.9)	(26.8)	8.8
Trade accounts payable	(9.5)	16.4	4.6
Other current and non-current assets	2.9	(15.3)	(1.6)
Other current and non-current liabilities	14.3	30.5	8.7
Current income taxes	(8.5)	(6.4)	10.2
EFFECT OF CHANGES IN ASSETS AND LIABILITIES ON OPERATING CASH FLOWS	(5.2)	(14.8)	(7.6)
NET CASH PROVIDED BY (USED IN) OPERATING ACTIVITIES	248.4	232.0	201.5
INVESTING ACTIVITIES			

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Investment in property, plant and equipment	(93.7)	(70.7)	(47.1)
Proceeds from sale of property, plant and equipment	1.1	0.1	0.1
Prepayments for other assets	-	-	(4.6)
Purchase of intangible assets	(0.5)	(1.2)	(0.5)
Acquisition of business, net of cash acquired	-	(35.0)	-
Sale of business, net of cash sold	11.5	-	-
NET CASH PROVIDED BY (USED IN) INVESTING ACTIVITIES	\$(81.6)	\$(106.8)	\$(52.1)

The accompanying Notes are an integral part of these financial statements.

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