
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 June 30, 2010

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 0-23125

OSI SYSTEMS, INC.

(Exact name of Registrant as specified in its charter)

Delaware
(State or Other Jurisdiction
of Incorporation or Organization)

12525 Chadron Avenue, Hawthorne, California
(Address of Principal Executive Offices)

Registrant's Telephone Number, Including Area Code: (310) 978-0516

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

Common Stock, \$0.001 par value

(Title of Class)

Securities registered pursuant to Section 12(g) of the Act: None

33-0238801
(I.R.S. Employer
Identification No.)

90250
(Zip 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 Website, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T during the preceding 12 months (or 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 the 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 definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act.

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:

The aggregate market value of the registrant's voting and non-voting Common Stock held by non-affiliates computed by reference to the price at which the Common Stock was last sold on December 31, 2009, the last business day of the registrant's most recently completed second fiscal quarter, was \$185,022,676.

The number of shares outstanding of the registrant's Common Stock as of August 24, 2010 was 18,404,606.

DOCUMENTS INCORPORATED BY REFERENCE

Portions of the definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders (to be filed subsequently) are incorporated by reference into Part III.

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PART I

Forward Looking Statements

This report contains “forward-looking statements” within the meaning of the Private Securities Litigation Reform Act of 1995, Section 27A of the Securities Act of 1933, as amended and Section 21E of the Securities Exchange Act of 1934, as amended. Forward-looking statements relate to expectations concerning matters that are not historical facts. Words such as “project,” “believe,” “anticipate,” “plan,” “expect,” “intend,” “may,” “should,” “will,” “would,” and similar words and expressions are intended to identify forward-looking statements. We believe that the expectations reflected in the forward-looking statements are reasonable, but those expectations may not prove to be correct. Important factors that could cause our actual results to differ materially from those expectations are disclosed in this report, including, without limitation, those described in Part I, Item 1, “Business,” Part I, Item 1A, “Risk Factors” and Part II, Item 7, “Management’s Discussion and Analysis of Financial Condition and Results of Operations” as well as elsewhere in this report and other documents previously filed or hereafter filed by us from time to time with the Securities and Exchange Commission. Such factors, of course, do not include all factors that might affect our business and financial condition. Although we believe that the assumptions upon which our forward-looking statements are based are reasonable, such assumptions could prove to be inaccurate and actual results could differ materially from those expressed in or implied by the forward-looking statements. All forward-looking statements contained in this report are qualified in their entirety by this statement. We undertake no obligation other than as may be required under securities laws to publicly update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

ITEM 1. BUSINESS

General

OSI Systems, Inc., together with its subsidiaries, is a vertically integrated designer and manufacturer of specialized electronic systems and components for critical applications. We sell our products in diversified markets, including homeland security, healthcare, defense and aerospace. Our company was originally incorporated in 1987 in California. In March 2010, we reincorporated our company in the State of Delaware. Our principal office is located at 12525 Chadron Avenue, Hawthorne, California 90250.

We have three operating divisions: (a) Security, providing security and inspection systems; (b) Healthcare, providing patient monitoring, diagnostic cardiology and anesthesia systems; and (c) Optoelectronics and Manufacturing, providing specialized electronic components for the Security and Healthcare divisions, as well as for applications in the defense and aerospace markets, among others.

Through our Security division, we design, manufacture and market security and inspection systems worldwide to end users under the “Rapiscan Systems” trade name. Rapiscan Systems products are used to inspect baggage, cargo, vehicles and other objects for weapons, explosives, drugs and other contraband, and to screen people. These products are also used for the safe, accurate and efficient verification of cargo manifests for the purpose of assessing duties and monitoring the export and import of controlled materials. Rapiscan Systems products fall into four categories: baggage and parcel inspection; cargo and vehicle inspection; hold (checked) baggage screening; and people screening. Our Security division also offers turnkey security screening services under the “S2 Global” trade name.

Through our Healthcare division, we design, manufacture and market patient monitoring, diagnostic cardiology and anesthesia delivery and ventilation systems worldwide to end users under the “Spacelabs” trade name. These products are used by care providers in critical care, emergency and perioperative areas within hospitals as well as physicians’ offices, medical clinics and ambulatory surgery centers.

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Through our Optoelectronics and Manufacturing division, we design, manufacture and market optoelectronic devices and provide electronics manufacturing services worldwide for use in a broad range of applications, including aerospace and defense electronics, security and inspection systems, medical imaging and diagnostics, computed tomography (CT), telecommunications and industrial automation. We sell our optoelectronic devices under the "OSI Optoelectronics" trade name and perform our electronics manufacturing services under the "OSI Electronics" trade name. We provide our optoelectronic devices and electronics manufacturing services to original equipment manufacturers, as well as to our own Security and Healthcare divisions.

In fiscal 2010, revenues from the Security division amounted to \$251.5 million, or approximately 42% of our revenues; revenues from the Healthcare division amounted to \$206.6 million, or approximately 35% of our revenues; and third-party revenues from the Optoelectronics and Manufacturing division amounted to \$137.1 million, or approximately 23% of revenues. Additional financial information concerning reporting segments and geographic areas is available in to our Consolidated Financial Statements, Note 15.

Industry Overview

We sell our security and inspection systems and patient monitoring, diagnostic cardiology and anesthesia systems primarily to end-users, while we design and manufacture our optoelectronic devices and value-added subsystems primarily for original equipment manufacturers.

Security. A variety of technologies are currently used worldwide in security and inspection applications, including transmission and backscatter X-ray, computed tomography, metal detection, trace detection, gamma-ray, passive millimeter wave, and neutron analysis. We believe that the market for security and inspection products will continue to be affected by the threat of terrorist incidents and by new government mandates and appropriations for security and inspection products in the United States and internationally.

The September 11, 2001 terrorist attacks on the World Trade Center and the Pentagon using hijacked airliners led to nationwide shifts in transportation and facilities security policies. Shortly following these attacks, Congress passed the Aviation and Transportation Security Act and integrated many U.S. security-related agencies, including the Federal Aviation Administration, into the U.S. Department of Homeland Security. Under its directive from Congress, the U.S. Department of Homeland Security has since undertaken numerous initiatives to prevent terrorists from entering the country, hijacking airliners, and obtaining and trafficking in weapons of mass destruction and their components, to secure sensitive U.S. technologies and to identify and screen high-risk cargo before it is loaded onto airlines and ships, among others. These initiatives, known, for example, as the Strategic Border Initiative, the Customs-Trade Partnership Against Terrorism and the U.S. Customs and Border Protection Container Security Initiative, have resulted in an increased demand for security and inspection products.

Recently, as part of the American Recovery and Reinvestment Act of 2009, Congress increased funding for a variety of U.S. Department of Homeland Security programs, including its Non-Intrusive Inspection (NII) Systems Program, which supports the deployment of security inspection systems used in the detection and prevention of contraband items such as weapons of mass effect, radioactive materials, narcotics, and currency, from entering or circulating within the United States.

Certain of the government sponsored initiatives in the United States, such as the U.S. Customs and Border Protection Container Security Initiative and the Customs-Trade Partnership Against Terrorism, have also stimulated security programs in other areas of the world because the U.S. initiatives call on other nations to bolster their port security strategies, including acquiring or improving their security and inspection equipment. The international market for non-intrusive inspection equipment, therefore, continues to expand as countries that ship goods directly to the United States participate in such programs and as they choose to procure equipment in order to secure their own borders, transportation networks, facilities and other venues.

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Congress also passed legislation that mandates the inspection of international maritime cargo destined for the United States, domestic civil aviation cargo, and for radiological and nuclear threats in cargo entering the United States. Certain of our cargo and vehicle inspection systems are already being used internationally and by the U.S. government to comply with these mandates.

Following recommendations outlined in the “9/11 Commission Report,” issued by the National Commission on Terrorist Attacks Upon the United States, the U.S. Department of Homeland Security now requires the screening of all cargo carried on passenger airlines in the United States. Several of our hold (checked) baggage and cargo screening systems have been approved by the U.S. Department of Homeland Security for this purpose and are being procured and used by freight forwarders, airlines, transportation companies and other businesses to fulfill their compliance requirements.

Following an attempted bombing on a Northwest airlines flight destined for Detroit, Michigan on Christmas Day 2009, on which a passenger tried to detonate explosives concealed beneath the passenger’s clothing, the U.S. Government initiated the widespread deployment of advanced imaging technology systems (body-scanners)—such as our Secure 1000 system—to U.S. airport checkpoints. These systems are used to detect both metallic and non-metallic threat objects concealed in or under clothing. This incident also prompted foreign governments to initiate similar deployments at other airports across the world.

Furthermore, the U.S. Department of Homeland Security’s Science and Technology Directorate has recently supported the development of new security inspection technologies and products. Our Security division participates in a number of such research and development efforts, including projects to develop new technologies for radiation and nuclear materials detection, aviation screening and suicide bomber detection. The Science and Technology Directorate has also initiated programs for the development of technologies capable of protecting highways, railways and waterways from terrorist attack.

In addition, the U.S. Department of Defense has begun to invest more heavily in technologies and services that screen would-be attackers before they are able to harm U.S. and allied forces. These technologies include products that can screen personnel, vehicles and other containers for the presence of explosives, improvised explosive devices (IEDs), weapons and other contraband.

Similar initiatives by international organizations such as the European Union have resulted in a growing worldwide demand for airline, cargo, port and border inspection technologies. For example, the European Union has issued uniform performance standards for systems that screen baggage and people at aviation checkpoints and air cargo, as well as new directives related specifically to maritime security, among others. Each of the performance standards is subject to ongoing revision as technological improvements occur in the marketplace. We anticipate that the promulgation of these new standards will continue to establish performance baselines against which our Security division will be able to direct certain of its research and development spending and market its products to customers located in the European Union.

As a result of these and other changes, sales of our security and inspection products have grown as compared to pre-September 11, 2001 levels. Major projects recently installed or currently underway include system installations at airports, ports and border crossings, government and military facilities and other locations in the United States and throughout the world. These projects contain various inspection product offerings. We anticipate that there may be growing demand from governments and commercial enterprises for increasingly sophisticated screening solutions in the future.

Healthcare. Healthcare has been, and we believe will continue to be, a growing sector throughout much of the world. Many developing countries in Asia and Latin America are expected to continue to build Healthcare infrastructure to serve expanding middle class populations. In developed countries, including the United States and Europe, an aging population is expected to fuel growth for many years.

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Many factors such as a nursing shortage in the United States and Europe, stricter government requirements affecting staffing and accountability as well as shrinking reimbursements from health insurance organizations are forcing healthcare providers to do more with less. Our Healthcare division designs, manufactures and markets products that respond to these economic forces by helping hospitals reduce costs while maintaining or improving the quality of care their physicians and nurses are able to deliver.

We are a global manufacturer and distributor of patient monitoring, cardiac monitoring and clinical networking solutions for use primarily in hospitals. We design, manufacture and market patient monitoring solutions for critical, emergency and perioperative care areas of the hospital, wired and wireless networks, and ambulatory blood pressure monitors, all aimed at providing caregivers with timely patient information. Our cardiac monitoring systems include Holter recorders, ECG, stress systems and related software and services. By making critical patient information more readily accessible both inside and outside the hospital, delays in decision-making can be reduced, length of stay can be shortened and treatment errors can be minimized.

We are also a global manufacturer and distributor of anesthesia delivery systems, ventilators and vaporizers. We sell these products primarily to hospitals for use in operating rooms and anesthesia induction areas as well as in magnetic resonance imaging (MRI) facilities. As pharmaceutical companies develop new anesthesia agents for the worldwide market, or as generic alternatives to patented anesthesia formulas become available, we work closely with them to support their new product introductions. As a result, we also sell systems and components, such as anesthesia vaporizers and ventilators, directly to pharmaceutical companies and other manufacturers of anesthesia delivery systems.

In October 2005, Spacelabs Healthcare, Inc., a subsidiary comprising the business operations of our Healthcare division, completed an initial public offering of approximately 20% of its total issued and outstanding common stock. The Spacelabs Healthcare shares traded under the ticker symbol "SLAB" on the AIM (formerly known as the Alternative Investment Market), a stock market administered by the London Stock Exchange. In the second quarter of fiscal 2007, we began repurchasing publicly-traded shares of Spacelabs Healthcare, increasing our ownership to 84% as of June 30, 2007. By December 31, 2007, we increased our ownership in Spacelabs Healthcare to 100% by repurchasing all remaining shares of Spacelabs Healthcare. Effective January 24, 2008, we cancelled Spacelabs Healthcare's AIM listing.

Optoelectronics and Manufacturing. Our optoelectronic devices are used in a wide variety of applications for diversified markets including the aerospace and defense, avionics, medical imaging and diagnostic, biochemistry analysis, pharmaceutical, nanotechnology, telecommunications, construction and homeland security markets. Medical applications for our devices include diagnostic and imaging products, computed tomography (CT) scanners, patient monitoring equipment, optometry instrumentation, and glucose monitors. Aerospace and defense applications for our devices include satellite navigation sensors, laser guided munitions systems, range finders, weapons simulation systems, computer peripherals and other applications that require the conversion of optical signals into electronic signals. Homeland security applications for our devices include X-ray-based and other detection systems. Our optoelectronic devices and value-added subsystems are also used in a wide variety of measurement control, monitoring and industrial applications and are key components in telecommunications technologies. We also offer electronics manufacturing services to our optoelectronics customers, as well as to our Security and Healthcare divisions. We have recently expanded such services by providing full turn-key and box-build manufacturing services, in which we provide product design and development, supply chain management, and production manufacturing services.

We believe that recent advances in technology and reductions in the cost of key components of optoelectronic systems, including computer processing power and memory, have broadened the market by enabling the use of optoelectronic devices in a greater number of applications. In addition, we see a trend among original equipment manufacturers to increasingly outsource the design and manufacture of optoelectronic devices as well as value-added subsystems to fully-integrated, independent manufacturers, like us, who may have greater specialization, broader expertise and the flexibility to respond in shorter time periods than most original

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equipment manufacturers can accomplish in-house. We believe that our level of vertical integration, substantial engineering resources, expertise in the use and application of optoelectronic technology and low-cost international manufacturing operations enable us to compete effectively in the market for optoelectronic devices and for electronics manufacturing services.

We have also penetrated several related markets that depend on our optoelectronic technologies and electronics manufacturing capabilities. For example, we sell a series of high-speed photodetectors for use in fiber optic systems such as gigabit ethernet, fiber channel and other telecommunication and data communication applications. Through system engineering and product development, we also develop, manufacture and sell laser-based remote sensing devices that are used to detect and classify vehicles in toll and traffic management systems and dual energy absorptiometry peripheral bone densitometers that are used to measure bone density in individuals that may be at risk for developing osteoporosis.

Growth Strategy

We believe that one of our primary competitive strengths is our expertise in the cost-effective design and manufacture of specialized electronic systems and components for critical applications. As a result, we have leveraged, and intend to continue to leverage, such expertise and capacity to gain price, performance and agility advantages over our competitors in the security, healthcare and optoelectronics fields, and to translate such advantages into profitable growth in those fields. At the same time, we continually seek to identify new markets in which our core expertise and capacity will provide us with competitive advantages. Key elements of this strategy include:

Capitalizing on Global Reach. We operate from locations in North America, Asia and Europe. We view our international operations as providing an important strategic advantage over competitors. First, international manufacturing facilities allow us to take advantage of competitive labor rates and favorable tax regulations in order to be a low cost producer. Second, our international offices strengthen our sales and marketing efforts and our ability to service and repair our systems by providing direct access to growing markets and to our existing international customer base. Third, multiple manufacturing locations allow us to reduce delivery times to our global customer base. In the future, we intend to develop new sources of manufacturing and sales capabilities to maintain and enhance the benefits of our international presence.

Capitalizing on Vertical Integration. Our vertical integration provides several advantages in each of our divisions. These advantages include reduced manufacturing and delivery times, lower costs due to our access to competitive international labor markets, direct sourcing of raw materials and quality control. We also believe that we offer significant added value to our customers by providing a full range of vertically-integrated services including component design and customization, subsystem concept design and application engineering, product prototyping and development, efficient pre-production and short-run and high volume manufacturing. We believe that our vertical integration differentiates us from many of our competitors and provides value to our customers who can rely on us to be an integrated supplier. We intend to continue to leverage our vertically integrated services to create greater value for our customers in the design and manufacture of our products.

Capitalizing on the Growing Market for Security and Inspection Systems. Heightened attentiveness to terrorist and other security threats may continue to drive growth in the market for security and inspection systems, not only in transportation security, but also at ports and border crossings, government installations, military facilities and public event venues. The trend toward increased screening of goods entering and departing from ports has resulted and many continue to result in growth in the market for cargo inspection systems and turnkey security screening services that are capable of screening shipping containers for contraband and assisting customs officials in the verification of shipping manifests. Package and cargo screening by freight forwarders, airlines and air cargo companies represents a growing sector, as new regulations in the U.S. and Europe require such screening in certain circumstances. In addition, the U.S. Congress recently concluded, as evidence by the American Recovery and Reinvestment Act of 2009, that public sector investment in security and inspection

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systems can help to stimulate growth in the U.S. economy. Following an attempted bombing on a Northwest airlines flight destined for Detroit, Michigan on Christmas Day 2009, on which a passenger tried to detonate explosives concealed beneath his clothing, the U.S. Government initiated the widespread deployment of advanced imaging technologies—such as the Secure 1000 body scanner that we offer—to screen passengers at aviation checkpoints. We intend to continue to expand our sales and marketing efforts both domestically and internationally, We further intend to capitalize on opportunities to replace, service and upgrade existing security installations as well as to offer turnkey security screening services in which we operate or assist in the operation of the installations. Finally, we also intend to continue to develop new security and inspection technologies, such as our real time tomography products, and to enhance our current product and services offerings through internal research and development and selective acquisitions in order to better address new applications and security industry demands.

Improving and Complementing Existing Medical Technologies. We develop and market patient monitoring systems, diagnostic cardiology products, anesthesia delivery systems, ventilators and vaporizers that utilize patient monitoring technologies. We are able to market and sell many of our product offerings through shared sales channels and distribution networks. Our efforts to develop new products and improve our existing medical technologies are focused on making decision-critical patient information available to care providers at the bedside, throughout a hospital, or even away from the hospital, thereby reducing time demands on physicians and nurses, enabling more rapid treatment decisions and improving patient care. Our efforts to improve existing diagnostic cardiology and anesthesia delivery technologies will also continue to concentrate on the development of devices that make it possible for institutions from large hospitals to small clinics and physicians' offices to obtain accurate, precise, reliable and cost-effective results.

Selectively Entering New Markets. We intend to continue to selectively enter new markets that complement our existing capabilities in the design, development and manufacture of specialized electronic systems and components for critical applications such as security and inspection and patient monitoring, diagnostic cardiology and anesthesia systems. We believe that by manufacturing end products that rely on our existing technological capabilities, we will leverage our integrated design and manufacturing infrastructure to capture greater margins and to build a larger presence in new end markets that present attractive competitive dynamics. We intend to achieve this strategy through internal growth and through selective acquisitions.

Acquiring New Technologies and Companies. Our success depends in part on our ability to continually enhance and broaden our product offerings in response to changing technologies, customer demands and competitive pressures. We have developed expertise in our various lines of business and other areas through internal research and development efforts as well as through selective acquisitions. As a vertically integrated designer and manufacturer of specialized electronic systems and components for critical applications, we have consistently looked for acquisition opportunities to broaden our technological expertise and capabilities, lower our manufacturing costs or facilitate our entry into new markets.

Products and Technology

We design, develop, manufacture and sell products ranging from security and inspection systems to patient monitoring, diagnostic cardiology and anesthesia systems to discrete optoelectronic devices and value-added subsystems.

Security and Inspection Systems. We design, manufacture and market security and inspection systems worldwide to end users under the “Rapiscan Systems” trade name. Rapiscan Systems products are used to inspect baggage, cargo, people, vehicles and other objects for weapons, explosives, drugs and other contraband. These systems are also used for the safe, accurate and efficient verification of cargo manifests for the purpose of assessing duties and monitoring the export and import of controlled materials. Rapiscan Systems products fall into four categories: baggage and parcel inspection, cargo and vehicle inspection, hold (checked) baggage screening and people screening. We also offer turnkey security screening services under the “S2 Global” trade name.

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As a result of the terrorist attacks of September 11, 2001, and subsequent attacks in other locations worldwide, security and inspection products have increasingly been used at a wide range of facilities other than airports, such as border crossings, railway stations, seaports, cruise line terminals, freight forwarding operations, government and military installations and nuclear facilities. As a result of the additional markets, we have successfully diversified our sales channels for security and inspection products.

Many of our security and inspection systems include dual- or multi-energy X-ray technology with computer software enhanced imaging technology to facilitate the detection of materials such as explosives, weapons, narcotics, currency or other contraband. While all X-ray systems produce a two-dimensional image of the contents of the inspected object, the dual-energy X-ray systems also measure the X-ray absorption of the inspected object's contents at two X-ray energies to determine the atomic number, mass and other characteristics of the object's contents. The various organic and inorganic substances in the inspected object appear to operators of the inspection systems in various colors and this visual information can be used to identify and differentiate the inspected materials. We have recently developed a proprietary dual-view X-ray technology, now available on many of our systems, that allows operators to examine objects from two orthogonal positions simultaneously, thereby reducing the need for re-scanning of objects and improving the operator's ability to detect threats. Our baggage and parcel inspection, cargo and vehicle inspection and hold baggage screening inspection systems range in size from compact tabletop systems to large systems comprising entire buildings in which trucks, shipping containers or pallets are inspected. Many of our inspection systems are also designed to be upgradeable to respond to new customer requirements as they emerge or change.

Our cargo and vehicle inspection applications, in which cars, trucks, shipping containers, pallets and other large objects can be inspected, are designed in various configurations, including fixed-site, gantry, relocatable, portal and mobile systems. These products are primarily used to verify the contents of cars, trucks or cargo containers and to detect the presence of contraband, including narcotics, weapons, explosives, and other smuggled items. They offer significant improvements over past methods of cargo screening, such as manual searches, as our cargo systems are faster, more thorough and do not subject the cargo to pilferage. Entire shipping containers or trucks containing densely packed goods can be screened rapidly.

Many of our cargo and vehicle inspection systems utilize X-ray or gamma-ray beams, in conjunction with digital imaging equipment, to non-intrusively inspect objects and present images to an inspector, showing shapes, sizes, locations and relative densities of the contents. Many of these systems have been built to meet specific customer inspection requirements.

Other cargo and vehicle inspection products automatically and non-intrusively detect chemical signatures indicating the presence of explosives and other contraband through the use of pulsed fast neutron and thermal neutron technologies, as opposed to ionizing radiation. Pulsed fast neutron and thermal neutron technologies permit the operator to inspect cargo, vehicles and containers based on the distinctive chemical composition of explosives, drugs or other contraband.

Our Security division is the only competitor in the market offering X-ray, gamma-ray and neutron-based material specific technologies. In addition, we are the only competitor in the marketplace that offers inspection systems at energy levels ranging from 200 KeV (Kilo electron Volts), to 1 MeV, 4.5 MeV, 6 MeV and 9MeV (Mega electron Volts). As a result, we believe that we offer the broadest technology platform in the cargo and vehicle inspection systems industry. This broad platform also permits us to offer customers hybrid solutions utilizing two or more of the technologies together, thereby optimizing flexibility, performance and cost to meet the customer's unique application requirements.

Our Security division also offers hold (checked) baggage screening systems that are utilized by airports, freight forwarders, and other parties responsible for screening baggage and cargo before it is placed in the cargo hold of airplanes. Our currently available systems utilize multiple, dual-energy X-ray beams to provide high-quality images to the operators of the systems and to enable detection algorithms that assist operators in the

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detection of explosives. These systems are designed to meet the high-speed screening and analysis demands of our customers. They can be operated in stand-alone mode, where a single operator views the images produced by a single system, or can be networked, allowing operators stationed at a remote computer terminal to monitor multiple systems.

Our Security division also offers people screening products such as a line of “Metor” brand walk-through metal detection products for use at security checkpoints at airports, amusement parks, banks, courthouses, government buildings, sports arenas and other venues, the WaveScan 200, which uses passive millimeter wave technology to screen for items concealed under clothing carried by individuals standing or walking at a distance, and the Secure 1000 personnel screener, which uses extremely low dose backscatter X-ray imaging to detect contraband and weapons concealed underneath clothing and hair. The Secure 1000 provides enhanced screening compared to metal detectors as it displays anomalies caused by very small amounts of metal as well as non-metallic items. As a result, the Secure 1000 can simultaneously locate and detect conventional metal weapons, as well as ceramic knives, explosives, illicit drugs, precious metals, cameras, recording devices and other contraband or security threats.

The following table sets forth certain information related to the standard security and inspection products that we currently offer. We do, however, also customize our standard products to suit specific applications and customer requirements.

<u>PRODUCT LINE</u>	<u>PRODUCT NAME / PRODUCT FAMILY</u>	<u>TECHNOLOGY</u>	<u>MARKET SEGMENT</u>
Baggage and Parcel Inspection	Rapiscan 500/600 series X-ray systems	Single and dual-energy X-ray	Checkpoint inspection at airports, prisons, border crossings, government buildings, postal facilities for mail screening
Cargo and Vehicle Inspection	Rapiscan Eagle Rapiscan VEDS Rapiscan GaRDS	High energy X-ray Thermal neutron analysis Gamma ray	Cargo and vehicle inspection at airports, border crossings and sea ports
Hold Baggage Screening	Rapiscan MVXR 5000	Multi-view, dual energy X-ray	Baggage inspection at airports and freight forwarding facilities
People Screening	Metor series metal detectors Rapiscan Secure 1000 WaveScan 200	Electromagnetic induction Backscatter X-ray Passive millimeter wave	Checkpoint inspection at airports, border crossings, stadiums, prisons and government facilities

Patient Monitoring, Diagnostic Cardiology and Anesthesia Systems. Our Healthcare businesses design, manufacture and market their products worldwide to end users primarily under the “Spacelabs” trade name.

Spacelabs products include patient monitors for use in perioperative, critical care and emergency care environments with neonatal, pediatric and adult patients. Our patient monitoring systems comprise monitors and central nursing stations connected via hardwired or wireless networks, as well as stand-alone monitors where the patient data can be transported physically from one monitor to another as the patient is moved. This ensures that hospital staff can access patient data where and when it is required. In addition, these products are designed with an “open architecture” to interact with hospital information systems. DNA, based on Citrix thin client technology, is a feature of many of these products which allows clinicians to view and control Microsoft Windows applications on the patient monitor’s display, eliminating the need for separate terminals in the patient’s room. Attending nurses can thereby check laboratory results and other reports, enter orders, review protocols and do charting right at the patient’s bedside. Inputs can be made using a mouse, keyboard and touchscreen.

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For electrocardiograph monitoring or multiparameter monitoring of ambulatory patients, we offer a digital telemetry system. The system operates in government-protected bands (608-614 MHz and 1.4GHz), not used for private land mobile radio, business radio services or broadcast analog and digital television. Our Ultraview digital telemetry solution comprises a lightweight and compact transmitter that enables monitoring of heart rate, ST segment, arrhythmia and continuous SpO₂ (pulse oximetry) monitoring. The multiparameter transmitter also integrates with our Ultralite ambulatory blood pressure monitor for the transmission of non-invasive blood pressure values to a central station or a multi-disclosure and information system.

In November 2009, we announced the expansion of our sales coverage in the United States by contracting with a manufacturer's representative organization to augment our direct sales channel. This addition of over 30 sales representatives enhances our ability to serve clinicians and healthcare facilities more effectively with Spacelabs' broad portfolio of patient monitors, telemetry, connectivity solutions, anesthesia systems, diagnostic cardiology products, clinician training and award-winning service. In addition, this new sales organization allows us to actively grow our expanding supplies and accessories business.

In April 2010, we launched the Ultraview DM3 dual-mode vital signs monitor. The Ultraview DM3 may function as either a spot check monitor or continuous function monitor, and provides caregivers with a dual-mode solution for accurately and efficiently measuring vital signs on adult and pediatric patients. Also in April 2010, we introduced the Ultraview SL2900 Dual Display monitor. The Ultraview SL2900 is designed to support the clinical requirements of the highest acuity environments. Dual displays allow a dedicated, full-screen view of real-time physiologic waveforms and alarms, plus an independent, full-screen presentation of charting and clinical applications.

We are also a leading supplier of ambulatory blood pressure monitors, which are routinely used in many European countries and are increasingly being used in the United States. Many physicians are using ambulatory blood pressure monitoring to detect "white coat" hypertension, a condition in which people experience elevated blood pressure in the doctor's office, but not in their daily lives. Ambulatory blood pressure monitoring is also used to adjust drug therapies for hypertensive patients. Ambulatory blood pressure monitoring helps improve diagnostic accuracy and minimize the associated costs of treatment.

Our Healthcare division develops cardiac monitoring systems, including Holter systems and recorders. Our Pathfinder and Impresario lines of Holter analyzers offer users interactive control with advanced diagnostic parameters. Our Lifecard and Aria recorders are worn by patients for up to seven days in order to capture heart arrhythmias that may occur in a patient only a few times per week. Patients that may be experiencing even less frequent heart arrhythmias wear our CardioCall product, which stays with the patient over several weeks and transmits its findings over the phone to a receiving station in the hospital. In addition to these products, we also offer other diagnostic cardiology products such as the CD12 electrocardiogram series and CH2000 stress test systems.

In May 2009, we launched our Sentinel product cardiology information management system. The Sentinel integrates data from Spacelabs-branded products into a central database that can be accessed by care providers and medical facility administrators. The Sentinel therefore provides enhanced workflow and efficiencies by centralizing recordings and reports into an enterprise wide system.

Our anesthesia delivery and ventilation group designs and manufactures anesthesia delivery systems, anesthesia vaporizers and ventilators. Our BleaseFocus, BleaseGenius and BleaseSirius anesthesia delivery systems provide flexible anesthesia solutions for most operating room environments, anesthesia induction areas, day surgery units, magnetic resonance imaging facilities and other areas where the administration of anesthesia is required. Our BleaseDatum anesthesia vaporizers and Blease 700/900 anesthesia ventilators are also designed to be compatible with the anesthesia delivery systems of several other manufacturers.

Recently, we added several new ventilators to our existing product line, each of which enables clinicians to enhance control over the delivery of ventilation and more finely tune their requirements to a surgical procedure

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and the individual characteristics of a patient. We added the BleaseSirius EFM product, still subject to FDA clearance, which allows the user to view information electronically rather than relying on traditional glass flow tubes, and provides for data transfer to an electronic patient medical record. Our new Manley anesthesia system incorporates the Blease 700/900 ventilator in a cost-effective, easy-to-use frame that allows high-end ventilation to care providers in emerging markets.

Additionally, we enhanced the product line with software updates that are intended to provide clinicians with greater ease of use as well as improvements and refinement of existing features and released an enhanced breathing system for the BleaseFocus and BleaseSerious product lines that allow clinicians to use lower gas flows, which is positive for the patient and the environment.

The following table sets forth a description of the more significant healthcare products that we currently offer:

<u>PRODUCT LINE</u>	<u>PRODUCT NAME / PRODUCT FAMILY</u>	<u>MARKET SEGMENT</u>
Patient Monitoring and Connectivity	Ultraview / Ultraview SL Intesys Clinical Suite G2 élançe MOM (Maternal Obstetrical Monitors)	All hospital care areas; outpatient surgery centers; and physician offices
Diagnostic Cardiology	Ambulatory blood pressure monitors Impresario Pathfinder CardioCall Lifecard Stress Testing Systems Sentinel ECG Data Management	All hospital cardiology care areas and physician offices
Anesthesia Delivery and Ventilation	Blease 700 and 900 series ventilators BleaseSirius BleaseSirius EFM BleaseDatum Vaporizer BleaseFocus BleaseGenius Manley Anesthesia System	Ambulatory surgery centers and operating rooms

Optoelectronic Devices and Manufacturing Services. Optoelectronic devices generally consist of both active and passive components. Active components sense light of varying wavelengths and convert the light detected into electronic signals, whereas passive components amplify, separate or reflect light. The active components we manufacture consist of silicon, gallium arsenide and indium gallium arsenide photodetectors. Passive components include lenses, prisms, filters, mirrors and other precision optical products that are used by us in the manufacture of our optoelectronic products or are sold to others for use in telescopes, laser printers, copiers, microscopes and other detection and vision equipment. The devices we manufacture are both standard products and products customized for specific applications and are offered either as components or as subsystems. For example, we have developed two-dimensional back-illuminated detector technology for security, medical and industrial computed tomography (CT) applications. This technology overcomes the limitations of conventional detectors by providing finer detector pitch density. This is used in high-resolution multi-slice CT scanners and other applications requiring improved image resolution. Our optoelectronic products and services are provided primarily under the “OSI Optoelectronics” trade name.

In addition to the manufacture of standard and original equipment manufacturer products, we also specialize in designing and manufacturing customized value-added subsystems for use in a wide range of products and

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equipment. An optoelectronic subsystem typically consists of one or more optoelectronic devices that are combined with other electronic components and packaging for use in an end product. The composition of a subsystem can range from a simple assembly of various optoelectronic devices that are incorporated into other subsystems (for example, a printed circuit board containing our optoelectronic devices) to complete end-products (for example, pulse oximetry equipment).

We have also expanded our electronics design and manufacturing capabilities both in North America and in Asia with enhanced, RoHS-compliant, printed circuit board and cable and harness manufacturing capabilities and box-build manufacturing services utilizing state-of-the-art automated surface mount technology lines. As a result, we now offer electronics manufacturing services for data and signal processing, amplifier and processor boards for medical equipment, musical tuning and studio hardware, motor controls, power supplies, and several other industrial applications that do not utilize optoelectronic devices. Our electronics manufacturing services are provided primarily under the “OSI Electronics” trade name.

We also develop, manufacture and sell laser-based remote sensing devices that are used to detect and classify vehicles in toll and traffic management systems under the “OSI Laserscan” trade name and we develop, manufacture and sell under the “Oseteometer” trade name, dual energy absorptiometry peripheral bone densitometers that are used to measure bone density in individuals that may be at risk for developing osteoporosis.

Markets, Customers and Applications

Security and Inspection Products. Most security and inspection products were developed in response to civilian airline hijackings. Consequently, a significant portion of our security and inspection products have been and continue to be sold for use at airports. However, our security and inspection products are also used for security purposes at locations in addition to airports, such as courthouses, office buildings, mailrooms, schools, freight forwarding facilities, prisons, high-profile locations such as Buckingham Palace, the Kremlin and the Vatican and for high-profile events such as the Olympic Games and World Cup Final. Furthermore, as terrorist attacks continue to occur, including recent attacks on civilian targets, overall transportation and travel industry demands have increased, resulting in heightened attention for our security and inspection products. In addition, our security and inspection products are increasingly being used for non-security purposes, such as for cargo inspection to detect narcotics and contraband and to verify manifests, prevention of pilferage at semiconductor manufacturing facilities, quality assurance and the detection of gold and currency.

Our customers include, among many others, the U.S. Transportation Security Administration, U.S. Customs and Border Protection, U.S. Department of Defense and Federal Bureau of Prisons, in the United States, as well as Her Majesty’s Revenue and Customs and Manchester Airport Group in the United Kingdom, Chek Lap Kok Airport in Hong Kong, Ben Gurion International Airport in Israel and the Malaysian Airport Board in Malaysia.

Patient Monitoring, Diagnostic Cardiology and Anesthesia Systems. Our patient monitoring, diagnostic cardiology and anesthesia systems are manufactured and distributed globally for use in critical care, emergency and perioperative areas within hospitals as well as physicians’ offices, medical clinics and ambulatory surgery centers. We also provide wired and wireless networks and clinical information access solutions, ambulatory blood pressure monitors and medical data services.

We have sold these products to organizations such as Eisenhower Medical Center in Rancho Mirage, California, Cape Fear Valley Health Systems in Fayetteville, North Carolina, Spartanburg Regional Medical Center in Spartanburg, South Carolina, LSU Medical Center in Shreveport, Louisiana, Schüchtermannklinik in Germany, LKW Villach in Austria and Universitätsspital Zürich in Switzerland, among many other organizations, including Premier, Inc., a hospital and healthcare system alliance with approximately 1,500 affiliated hospitals and other healthcare sites.

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Optoelectronic Devices and Electronics Manufacturing Services. Our optoelectronic devices and the electronics we manufacture are used in a broad range of products by a variety of customers. For example, they are utilized by customers in the following market segments: aerospace and avionics; analytical and medical imaging; telecommunications; homeland security; healthcare; military defense; office automation; and toll and traffic management. Major customers in these segments include SCM, Honeywell, Flir Systems, Raytheon, JDS Uniphase, ITT Corporation, Gilardoni, Bayer, Covidien, Smiths Medical, Beckman Coulter, Lockheed Martin, United Technologies and Northrop Grumman, among others.

Marketing, Sales and Service

We market and sell our security and inspection products worldwide through a direct sales and marketing staff of approximately 100 employees located in North America, Europe, Asia and Australia, in addition to an expansive global network of independent distributors. This sales staff is supported by a service organization located primarily in North America, Europe and Asia, as well as a global network of independent distributors. We also support these sales and customer relations efforts by providing operator training, computerized training and testing equipment, in-country service support, software upgrades and service training for customer technicians.

We market and sell our patient monitoring, diagnostic cardiology and anesthesia systems worldwide through a direct sales and marketing staff of approximately 230 sales personnel and 190 service personnel located in North America, Europe and Asia, in addition to a global network of independent distributors. We also support these sales and customer service efforts by providing operator in-service training, software updates and upgrades and service training for customer biomedical staff and distributors.

We market and sell our optoelectronic devices and value-added manufacturing services, through both a direct sales and marketing staff of approximately 40 employees located in North America, Europe and Asia, and indirectly through a global network of independent sales representatives and distributors. Our sales staff is supported by an applications engineering group whose members are available to provide technical support, which includes designing applications, providing custom tooling and process integration and developing products that meet customer defined specifications.

We consider our maintenance service operations to be an important element of our business. After the expiration of our standard product warranty periods, we are sometimes engaged by our customers to provide maintenance services for our security and inspection products through annual maintenance contracts. We provide a variety of service and support options for our patient monitoring, diagnostic cardiology and anesthesia systems customers, ranging from complete on-site repair and maintenance service and telephone support to parts exchange programs for customers with the internal expertise to perform a portion of their own service needs. We believe that our international maintenance service capabilities allow us to be competitive in selling our security and inspection systems as well as our patient monitoring, diagnostic cardiology and anesthesia systems. Furthermore, we believe that as the installed base of both our security and inspection systems and patient monitoring, diagnostic cardiology and anesthesia systems increases, revenues generated from such annual maintenance service contracts and from the sale of replacement parts will increase.

Research and Development

Our security and inspection systems are primarily designed at our facilities in the United States and internationally in Finland, Malaysia, India and the United Kingdom. These products include mechanical, electrical, analog electronic, digital electronic and software subsystems, which are all designed by us. In addition to product design, we provide system integration services to integrate our products into turnkey systems at the customer site. We support cooperative research projects with government agencies and provide contract research for government agencies.

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Our patient monitoring, diagnostic cardiology and anesthesia systems are primarily designed at our facilities in the United States and internationally in China, India and the United Kingdom. Such systems include mechanical, electrical, digital electronic and software subsystems, all of which are designed by us. We are also currently involved, both in the United States and internationally, in several research projects aimed at improving our medical systems and at expanding our current product line.

We design and manufacture optoelectronic devices and we provide electronics manufacturing services primarily in our facilities in the United States and internationally in India, Indonesia, Malaysia and Singapore. We engineer and manufacture subsystems to solve the specific application needs of our original equipment manufacturer customers. In addition, we offer entire subsystem design and manufacturing solutions. We consider our engineering personnel to be an important extension of our core sales and marketing efforts.

In addition to close collaboration with our customers in the design and development of our current products, we maintain an active program for the development and introduction of new products, enhancements and improvements to our existing products, including the implementation of new applications of our technology. We seek to further enhance our research and development program and consider such program to be an important element of our business and operations. As of June 30, 2010, we engaged approximately 397 full-time engineers, technicians and support staff. Our research and development expenses were \$45.3 million in fiscal 2008, \$36.9 million in fiscal 2009 and \$38.6 million in fiscal 2010. We intend to continue to invest in our research and development efforts in the future.

Manufacturing and Materials

We currently manufacture our security and inspection systems domestically in California, Mississippi and North Carolina, and internationally in Malaysia and the United Kingdom. We currently manufacture our patient monitoring, diagnostic cardiology and anesthesia systems domestically in Washington, and internationally in China. We currently manufacture our optoelectronic devices and provide electronics manufacturing services domestically in California, Massachusetts and Mississippi, and internationally in India, Indonesia, Malaysia, and Singapore. Most of our high volume, labor intensive manufacturing and assembly activities are performed at our facilities in India, Indonesia and Malaysia. Since most of our customers are located in the United States, Europe and Asia, our ability to manufacture products in these markets and provide follow-on service from offices located in these regions is an important component of our global strategy.

Our global manufacturing organization has expertise in optoelectronic, microelectronic and integrated electronics for commercial, medical, aerospace and defense industry applications. Our manufacturing includes silicon wafer processing and fabrication, optoelectronic device assembly and screening, thin and thick film microelectronic hybrid assemblies, surface mounted and thru-hole printed circuit board electronic assemblies and electronics services, including complete turn-key and box-build manufacturing. We outsource certain manufacturing operations, including certain sheet metal fabrication and plastic components. The manufacturing process for components and subsystems consists of manual tasks performed by skilled technicians as well as automated tasks.

The principal raw materials and subcomponents used in producing our security and inspection systems consist primarily of X-ray generators, linear accelerators, radioactive isotopes, neutron generators, detectors, data acquisition and computer systems, and conveyance systems. A large portion of the optoelectronic devices, subsystems and circuit card assemblies used in our inspection and detection systems are manufactured in-house. The metal enclosures used in our baggage and parcel inspection systems are also manufactured in-house, while the X-ray generators, linear accelerators, radioactive isotopes, neutron generators and conveyance systems used in our cargo and vehicle inspection systems are purchased from unaffiliated third party providers.

The principal raw materials and subcomponents used in producing our patient monitoring, diagnostic cardiology and anesthesia systems consist of printed circuit boards, housings, mechanical assemblies, pneumatic

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devices, cables, filters and packaging materials. We purchase certain devices, including computers, peripheral accessories and remote displays from unaffiliated third party providers.

The principal raw materials and subcomponents used in producing our optoelectronic devices and electronic subsystems consist of silicon wafers, electronic components, light emitting diodes, scintillation crystals, passive optical components, printed circuit boards, and packaging materials. The silicon-based optoelectronic devices manufactured by us are critical components in most of our products and subsystems. We purchase silicon wafers and other electronic components from unaffiliated third party providers.

For cost, quality control and efficiency reasons, at times we purchase raw materials and subcomponents only from single vendors with whom we have ongoing relationships. We do, however, qualify second sources for most of our raw materials and critical components. We purchase the materials pursuant to purchase orders placed from time to time in the ordinary course of business. Although to date none of our divisions has experienced any significant shortages or material delays in obtaining any of its raw materials or subcomponents, it is possible that they may face such shortages or delays in one or more materials in the future.

Patents, Trademarks, Tradenames and Licenses

Trademarks and Tradenames. We have used, registered and applied to register certain trademarks and service marks to distinguish our products, technologies and services from those of our competitors in the United States and in foreign countries. We enforce our trademark, service mark and trade name rights in the United States and abroad.

Patents. We hold a number of U.S. and foreign patents relating to various aspects of our security and inspection products, patient monitoring, diagnostic cardiology and anesthesia systems and optoelectronic devices and subsystems. Our current patents will expire at various times between 2010 and 2030. However, it remains possible that pending patent applications or other applications that may be filed may not result in issued patents. In addition, issued patents may not survive challenges to their validity. Although we believe that our patents have value, our patents, or any additional patents that may be issued in the future, may not be able to provide meaningful protection from competition.

Licenses. Our Security, Healthcare and Optoelectronics and Manufacturing divisions have each entered into a variety of license arrangements under which they are permitted to manufacture, market, and/or sell a limited number of the products that we offer and/or to service various types of software, data, equipment, components and enhancements to our own proprietary technology.

We believe that our trademarks and tradenames, patents and licenses are important to our business. The loss of some of our trademarks, patents or licenses might have a negative impact on our financial results and operations. However, we operate in a competitive environment with a known customer base and rely mainly on providing our customers with quality products and services to ensure continuing business. Thus, with the exception of the loss of either the Spacelabs® or Rapiscan® trademarks, the impact of the loss of any single trademark, patent or license would not likely have a material adverse effect on our business. We consider the Spacelabs® trademark an important asset and have registered it in approximately forty countries. In addition, we have instituted a similar registration program for the Rapiscan® trademark.

Regulation of Medical Products

The patient monitoring, diagnostic cardiology and anesthesia systems we manufacture and market are subject to regulation by numerous government agencies, principally the U.S. Food and Drug Administration (FDA) and by certain state and foreign authorities. They are also subject to various U.S. and foreign electrical safety standards.

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The FDA has broad regulatory powers with respect to pre-clinical and clinical testing of new medical products and the designing, manufacturing, marketing and advertising of medical products. It requires that all medical devices introduced into the market be preceded either by a pre-market notification clearance order under section 510(k) of the Food, Drug and Cosmetic Act, or an approved pre-market approval application. A 510(k) pre-market notification clearance order indicates that the FDA agrees with an applicant's determination that the product for which clearance has been sought is substantially equivalent to another legally marketed medical device. The clearance of a pre-market approval application, on the other hand, indicates that the FDA has determined that the device has been proven, through the submission of clinical trial data and manufacturing quality assurance information, to be safe and effective for its labeled indications. The process of obtaining 510(k) clearance typically takes between three and six months, but can take substantially longer. The pre-market approval application review process, on the other hand, can last more than a year. To date, all of the patient monitoring, diagnostic cardiology and anesthesia systems we manufacture and sell in the United States have required only 510(k) pre-market notification clearance.

Such regulatory approvals, when granted, may entail limitations on the indicated uses for which a product may be marketed, and such product approvals, once granted, may be withdrawn if problems occur after initial marketing. Manufacturers of FDA-regulated products are subject to pervasive and continuing governmental regulation, including extensive recordkeeping requirements and reporting of adverse experiences associated with product manufacture and use. Compliance with these requirements is costly, and failure to comply can result in, among other things, fines, total or partial suspension of production, product recalls, failure of the FDA to review pending marketing clearances or approval applications, withdrawal of marketing clearances or approvals or even criminal prosecution.

We are also subject to regulation in the foreign countries in which we manufacture and market our patient monitoring, diagnostic cardiology and anesthesia systems. For example, the commercialization of medical devices in the European Union is regulated under a system that presently requires all medical devices sold in the European Union to bear the CE mark—an international symbol of adherence to quality assurance standards. Our manufacturing facilities in Hawthorne, California; Issaquah, Washington; and Suzhou in China are all certified to the International Organization for Standardization's ISO 13485 standard for medical device quality management systems. The Hawthorne, California and Issaquah, Washington facilities are also certified to the requirements of Annex II, section 3 of the Directive 93/42 EEC on Medical Devices, which allows them to self-certify that newly manufactured products can bear the CE mark.

We believe we are in compliance with all applicable federal, state and foreign regulations regarding the manufacture and sale of our patient monitoring, diagnostic cardiology and anesthesia delivery systems except to an extent that would not have a material adverse effect on our business, financial condition or results of operations. Such regulations and their enforcement do, however, constantly change, and we cannot predict what effect, if any, such changes may have on our businesses in the future.

Environmental Regulations

We are subject to various federal, state and local environmental laws, ordinances and regulations relating to the use, storage, handling and disposal of certain hazardous substances and wastes used or generated in the manufacturing and assembly of our products. Under such laws, we may become liable for the costs of removal or remediation of certain hazardous substances that have been released on or in our facilities or that have been disposed of off-site as waste. Such laws may impose liability without regard to whether we knew of, or caused, the release of such hazardous substances. We have conducted Phase I environmental site assessments for each of our properties in the United States at which we manufacture products. The purpose of each such report is to identify, as of the date of such report, potential areas of environmental concern related to past and present activities or from nearby operations. In certain cases, we have conducted further environmental assessments consisting of soil and groundwater testing and other investigations deemed appropriate by independent environmental consultants. We believe that, except to an extent that would not have a material adverse effect on

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our business, financial condition or results of operations, we are currently in compliance with all environmental regulations in connection with our manufacturing operations, and that we have obtained all environmental permits necessary to conduct our business. The amount of hazardous substances and wastes produced and generated by us may increase in the future depending on changes in our operations. Any failure by us to comply with present or future regulations could subject us to the imposition of substantial fines, suspension of production, alteration of manufacturing process or cessation of operations, any of which could have a material adverse effect on our business, financial condition and results of operations.

During one investigation, we discovered soil and groundwater contamination at our Hawthorne, California facility. We filed the requisite reports concerning this problem with the appropriate environmental authorities in fiscal 2001. We have not yet received any response to such reports, and no agency action or litigation is presently pending or threatened. We also have notified the prior owners of the facility and the present owners and tenants of adjacent properties concerning the problem and have requested from such parties agreements to toll of the statute of limitations with respect to actions against such parties with respect to the contamination in order that we may focus our attention on resolution of the contamination problem. Our site was previously used by other companies for semiconductor manufacturing similar to that presently conducted on the site by us, and it is not presently known who is responsible for the contamination or, if required, the remediation. The groundwater contamination is a known regional problem, not limited to our premises or our immediate surroundings.

We have also been informed of soil and groundwater evaluation efforts at a facility that our Ferson Technologies subsidiary previously leased in Ocean Springs, Mississippi. Ferson Technologies occupied the facility between 1993 and 2003. We believe that the owner and previous occupants of the facility have primary responsibility for any remediation that may be required and have an agreement with the facility's owner under which the owner is responsible for remediation of pre-existing conditions. However, as site evaluation efforts are still in progress, and may be for some time, we are unable at this time to ascertain whether Ferson Technologies bears any exposure for remediation costs under applicable environmental regulations.

Competition

The markets in which we operate are highly competitive and characterized by evolving customer needs and rapid technological change. We compete with a number of other manufacturers, some of which have significantly greater financial, technical and marketing resources than we have. In addition, these competitors may have the ability to respond more quickly to new or emerging technologies, adapt more quickly to changes in customer requirements, have stronger customer relationships, have greater name recognition and may devote greater resources to the development, promotion and sale of their products than we do. As a result, we may not be able to compete successfully against designers and manufacturers of specialized electronic systems and components, broadly speaking, or more specifically within the markets for security and inspection systems, patient monitoring, diagnostic cardiology and anesthesia systems, or optoelectronic devices. Future competitive pressures may materially and adversely affect our business, financial conditions and results of operations.

In the security and inspection market, competition is based primarily on such factors as product performance, functionality and quality, the overall cost effectiveness of the system, prior customer relationships, technological capabilities of the products, price, local market presence and breadth of sales and service organization. We believe that our principal competitors in the market for security and inspection products are Smiths Detection; L-3 Communications—Security and Detection Systems division; American Science and Engineering; Morpho Detection; SAIC; CEIA and Nuctech. Competition could result in price reductions, reduced margins and loss of market share. We also compete for potential customers based on existing relationships between our competitors and the customers. Although we also have established relationships with a number customers, we may not be able to compete successfully in the future with existing competitors or new entrants. In the cargo and vehicle inspection systems market, we compete for potential customers based on price, performance and the ability to design both standard and customized products. Several of our competitors have operated in this area for longer than we have. However, due to our recent successes in designing and delivering

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high-energy X-ray and gamma-ray systems, we believe that we have demonstrated an ability to compete effectively. Additionally, although our competitors in the cargo and vehicle inspection market each offer products in competition with one or more of our products, our ability to supply a variety of high-energy X-ray and gamma-ray systems means that we offer among the widest array of solutions available from a single supplier. This variety of technologies also permits us to offer unique hybrid systems to our customers that utilize two or more of these technologies, thereby optimizing flexibility, performance and cost to meet the customer's unique application requirements.

In the patient monitoring, diagnostic cardiology and anesthesia systems delivery market, competition is also based on a variety of factors including product performance, functionality, value and breadth of sales and service organization. We believe that our principal competitors in the market for patient monitoring, diagnostic cardiology and anesthesia systems are Philips Medical; GE Healthcare; Mindray Medical, Cardiac Science; Mortara Instrument; Dräger Medical; Nihon Kohden; Penlon and Marquet. Competition could result in price reductions, reduced margins and loss of our market share. We believe that our patient monitoring products are easier to use than the products of many of our competitors because we offer a consistent user interface throughout many of our product lines. Finally, while some of our competitors are also beginning to introduce portal technology, which allows remote access to data from the bedside monitor, central station or other point of care, we believe that our competing technologies are superior in bringing instant access to labs, radiology and charting at the point of care. Although we have established relationships with a number of large hospitals, we may not be able to successfully compete in the future with existing competitors or with new entrants.

In the markets in which we compete to provide optoelectronic devices and electronics manufacturing services, competition is based primarily on such factors as expertise in the design and development of optoelectronic devices, product quality, timeliness of delivery, price, customer technical support and on the ability to provide fully integrated services from application development and design through production. We believe that our major competitors in the optoelectronic device market are PerkinElmer and Hamamatsu. Because we specialize in custom subsystems requiring a high degree of engineering expertise, we believe that we generally do not compete to any significant degree with any other large United States, European or Asian manufacturers of standard optoelectronic components. Competition in the extensive electronic manufacturing services market ranges from multinational corporations with sales in excess of several billions of dollars, to large regional competitors and to small local assembly companies. In our experience, the original equipment manufacturers to whom we provide such services prefer to engage companies that offer both local and lower-cost off-shore facilities. As a result, our primary domestic competition for these services is located in Southern California and in New England, where our U.S. facilities are also located. Such competition includes CTS; Stellar Microelectronics; Senior Systems Technology; Celestica and Benchmark Electronics, among others. In addition, our high-volume, low-cost contract manufacturing locations in Southeast Asia compete with other manufacturers in the same region.

Backlog

We measure our backlog as orders for which purchase orders or contracts have been signed, but which have not yet been shipped and for which revenues have not yet been recognized.

We ship most of our baggage and parcel inspection, hold (checked) baggage screening, people screening, patient monitoring, diagnostic cardiology and anesthesia systems and optoelectronic devices and value-added subsystems within one to several months after receiving an order. However, such shipments may be delayed for a variety of reasons, including any special design or requirements of the customer. In addition, large orders of security and inspection products typically require greater lead-times.

Certain of our cargo and vehicle inspection and hold (checked) baggage screening systems may require several months lead-time. We have experienced some significant shipping delays associated with our cargo and vehicle inspection systems. Such delays can occur for many reasons, including: (i) additional time necessary to

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conduct inspections at the factory before shipment; (ii) a customer's need to engage in time-consuming special site preparation to accommodate the system, over which we have no control or responsibility; (iii) additional fine tuning of such systems once they are installed; (iv) design or specification changes by the customer; and (v) delays originating from other contractors on the project.

As of June 30, 2010, our consolidated backlog totaled approximately \$240 million, compared to approximately \$203 million as of June 30, 2009 and approximately \$212 million at June 30, 2008. Sales orders underlying our backlog are firm orders; although, from time to time we may agree to permit a customer to cancel an order or an order may be cancelled due to protest. Variations in the size of orders, product mix, or delivery requirements, among other factors, may result in substantial fluctuations in backlog from period to period. Backlog as of any particular date should not be relied upon as indicative of our revenues for any future period and cannot be considered a meaningful indicator of our performance on an annual or quarterly basis.

Employees

As of June 30, 2010, we employed approximately 3,183 people, of whom 1,616 were employed in manufacturing, 397 were employed in engineering or research and development, 385 were employed in administration, 367 were employed in sales and marketing and 418 were employed in service capacities. Of the total employees, approximately 1,391 were employed in North America and South America, 1,405 were employed in Asia and 387 were employed in Europe. Many of our employees in Europe have statutory collective bargaining rights. We have never experienced a work stoppage or strike, and management believes that its relations with employees are good.

Available Information

We are subject to the informational requirements of the Securities Exchange Act of 1934, as amended. Therefore, we file periodic reports, proxy statements and other information with the Securities and Exchange Commission. Such reports, proxy statements and other information may be obtained by visiting the Public Reference Room of the Securities and Exchange Commission at 100 F Street, N.E., Washington, D.C. 20549 or by calling the Securities and Exchange Commission at 1-800-SEC-0330. In addition, the Securities and Exchange Commission maintains an Internet website (<http://www.sec.gov>) that contains reports, proxy statements and other information that issuers are required to file electronically.

Our Internet address is: <http://www.osi-systems.com>. We make available, free-of-charge through our Internet website, our annual report on Form 10-K, quarterly reports on Form 10-Q, current reports on Form 8-K, amendments to those reports filed or furnished pursuant to Section 13(a) or 15(d) of the Securities Exchange Act of 1934, as amended and reports filed pursuant to Section 16 of the Securities Exchange Act of 1934, as amended. We do so as soon as reasonably practicable after electronically filing such material with, or furnishing it to, the Securities and Exchange Commission.

ITEM 1A. RISK FACTORS

We encourage you to carefully consider all of the following risk factors when making investment decisions regarding our company. If any of the following risks materialize, our business, financial condition and operating results could be materially adversely affected.

Fluctuations in our operating results may cause our stock price to decline.

Given the nature of the markets in which we participate, it is difficult to reliably predict future revenues and profitability. Changes in competitive, market and economic conditions may cause us to adjust our operations. A high proportion of our costs are fixed, due in part to our significant sales, research and development and

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manufacturing costs. Thus, small declines in revenue could disproportionately affect our operating results. Factors that may affect our operating results and the market price of our Common Stock include:

- demand for and market acceptance of our products;
- competitive pressures resulting in lower selling prices;
- adverse changes in the level of economic activity in regions in which we do business;
- low or fluctuating levels of political stability in regions in which we do business;
- adverse changes in industries, such as semiconductors and electronics, on which we are particularly dependent;
- changes in the portions of our revenue represented by various products and customers;
- delays or problems in the introduction of new products;
- the announcement or introduction of new products, services or technological innovations by our competitors;
- variations in our product mix;
- the timing and amount of our expenditures in anticipation of future sales;
- the availability of equity and credit markets to provide our customers with funding to make equipment purchases;
- exchange rate fluctuations;
- increased costs of raw materials or supplies;
- changes in the volume or timing of product orders;
- timing of completion of acceptance testing of some of our products;
- changes in regulatory requirements;
- natural disasters; and
- changes in general economic factors.

Unfavorable currency exchange rate fluctuations could adversely affect our profitability.

Our international sales and our operations in foreign countries expose us to risks associated with fluctuating currency values and exchange rates. Gains and losses on the conversion of accounts receivable, accounts payable and other monetary assets and liabilities to U.S. dollars may contribute to fluctuations in our results of operations. In addition, increases or decreases in the value of the U.S. dollar relative to other currencies could have an adverse effect on our results of operations.

We face aggressive competition in many areas of business. If we do not compete effectively, our business will be harmed.

We encounter aggressive competition from numerous competitors in many areas of our business. In the security and inspection and patient monitoring, diagnostic cardiology and anesthesia systems markets, competition is based primarily on such factors as product performance, functionality and quality, cost, prior customer relationships, technological capabilities of the product, price, certification by government authorities, local market presence and breadth of sales and service organization. In the optoelectronic devices and electronics manufacturing markets, competition is based primarily on factors such as expertise in the design and development of optoelectronic devices, product quality, timeliness of delivery, price, customer technical support and on the ability to provide fully-integrated services from application development and design through volume

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subsystem production. We may not be able to compete effectively with all of our competitors. To remain competitive, we must develop new products and enhance our existing products and services in a timely manner. We anticipate that we may have to adjust prices of many of our products to stay competitive. In addition, new competitors may emerge, and entire product lines or service offerings may be threatened by new technologies or market trends that reduce the value of these product lines or service offerings.

The September 11, 2001 terrorist attacks and the creation of the U.S. Department of Homeland Security have increased financial expectations that may not materialize.

The September 11, 2001 terrorist attacks and the subsequent creation of the U.S. Department of Homeland Security have created increased interest in our security and inspection systems. However, we are not certain whether the level of demand will continue to be as high as it is now. We do not know what solutions will continue to be adopted by the U.S. Department of Homeland Security, the U.S. Department of Defense, and similar agencies in other countries and whether our products will be a part of those solutions. Additionally, should our products be considered as a part of the future security solutions, it is unclear what the level may be and how quickly funding to purchase our products may be made available. These factors may adversely impact us and create unpredictability in revenues and operating results.

If operators of, or algorithms installed in, our security and inspection systems fail to detect weapons, explosives or other devices that are used to commit a terrorist act, we could be exposed to product and professional liability and related claims for which we may not have adequate insurance coverage.

Our business exposes us to potential product liability risks that are inherent in the development, manufacturing, sale and service of security inspection systems as well as in the provision of training to our customers in the use and operation of such systems. Our customers use our security and inspection systems to help them detect items that could be used in performing terrorist acts or other crimes. Some of our security and inspection systems require that an operator interpret an image of suspicious items within a bag, parcel, container or other vessel. Others signal to the operator that further investigation is required. In either case, the training, reliability and competence of the customer's operator are crucial to the detection of suspicious items.

Security inspection systems that signal to the operator that further investigation is required are sometimes referred to in the security industry as "automatic" detection systems. Such systems utilize software algorithms (often designed to meet government requirements) to interpret data produced by the system and to signal to the operator when a dangerous object may be present. Such algorithms are probabilistic in nature and are also subject to significant technical limitations. Nevertheless, if such a system were to fail to signal to an operator when an explosive or other contraband was in fact present, resulting in significant damage, we could become the subject of significant product liability claims.

Furthermore, security inspection by technological means is always circumstance and application-specific. In addition, our security and inspection systems are not designed to work under all circumstances. We test the reliability of our security and inspection systems during both their development and manufacturing phases. We also perform such tests if we are requested to perform installation, warranty or post-warranty servicing. However, our security inspection systems are advanced mechanical and electronic devices and therefore can malfunction.

We also offer turnkey security screening services under which we perform certain of the security screening tasks that have historically been performed by our customers. Such tasks include: design and layout of the security checkpoint where the inspection equipment is located; selection of the security equipment to be used at the checkpoint; selection, training and management of the personnel operating the checkpoint; operation of the security screening equipment itself; interpretation of the images and other signals produced by the security screening equipment; maintenance and security of the checkpoint as well as other related services. Such projects expose us to certain professional liability risks that are inherent in performing security inspection services (in live checkpoint environments and over an extended period of time) for the purpose of assisting our customers in the

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detection of contraband items, including items that could be used in performing terrorist acts or other crimes. Although we utilize operational and other procedures, many of which are designed to our customers' specifications and requirements, intended to limit the risk of failure, if a contraband item passes through the checkpoint and is used to perform a terrorist act or other crime, we could become the subject of significant professional liability claims.

In addition, there are also many other factors beyond our control that could lead to liability claims should an act of terrorism occur. The September 11, 2001 and 1993 World Trade Center bombing attacks, and the potential for future attacks, have caused commercial insurance for such threats to become extremely difficult to obtain. It is very likely that, should we be found liable following a major act of terrorism, the insurance we currently have in place would not fully cover the claims for damages.

Our patient monitoring, diagnostic cardiology and anesthesia systems could give rise to product liability claims that could materially and adversely affect our financial condition and results of operations.

The development, manufacturing and sale of medical devices expose us to significant risk of product liability claims and, sometimes, product failure claims. We face an inherent business risk of financial exposure to product liability claims if the use of our medical devices results in personal injury or death. Substantial product liability litigation currently exists within the medical device industry. Some of our patient monitoring, diagnostic cardiology and anesthesia systems businesses are, from time to time, subject to product liability claims and/or product recalls. Future product liability claims may exceed the limits of our insurance coverages or such insurance may not continue to be available to us on commercially reasonable terms, or at all. Consequently, a product liability claim or other claim with respect to uninsured liabilities, or in excess of insured liabilities, could have a material adverse effect on our business, financial condition, operating results and cash flows.

If we are unable to sustain high quality processes for the manufacture and delivery of goods and services, our reputation could be harmed, our competitive advantage could erode and we could incur significant costs.

Quality is extremely important to us and our customers due in part to the serious consequences of product failure. Our quality certifications are critical both to the marketing success of our goods and services and to the satisfaction of both regulatory and contractual requirements under which we sell many of our products. If we fail to meet these standards or other standards required in our industries, we could lose customers and market share, our revenue could decline and we could face significant costs and other liabilities.

The loss of certain of our customers could have a negative effect on our reputation.

We sell many of our products to prominent, well-respected institutions, including agencies and departments of the U.S. Government, state and local governments, foreign governments, renowned hospitals and hospital networks, and large military-defense and space-industry contractors. Many of these larger customers spend considerable resources testing and evaluating our products and our design and manufacturing processes and services. Some of our smaller customers know this and rely on this as indication of the high-quality and reliability of our products and services. As a result, part of our reputation and success depends on our ability to continue to sell to larger institutions that are known for demanding high standards of excellence. The loss or termination of a contract by such an institution, even if for reasons unrelated to the quality of our products or services, could therefore have a more wide-spread and potentially material adverse effect on our business, financial condition and results of operations.

Our revenues are dependent on orders of security and inspection systems and patient monitoring, diagnostic cardiology and anesthesia systems, which may have lengthy and unpredictable sales cycles.

Sales of security and inspection systems often depend upon the decision of governmental agencies to upgrade or expand existing airports, border crossing inspection sites, seaport inspection sites and other security installations. Sales outside of the United States of our patient monitoring, diagnostic cardiology and anesthesia

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systems depend in significant part on the decision of governmental agencies to build new medical facilities or to expand or update existing medical facilities. Accordingly, a significant portion of our sales of security and inspection systems and our patient monitoring, diagnostic cardiology and anesthesia systems is often subject to delays associated with the lengthy approval processes that typically accompany such capital expenditures. During these approval periods, we expend significant financial and management resources in anticipation of future orders that may not occur. If we fail to receive an order after expending such resources, such failure could have a material adverse effect on our business, financial condition and results of operations.

Current economic conditions, including the slow pace of recovery from recession in the United States and other parts of the world, as well as further disruptions in the financial markets could result in substantial declines in our revenues, earnings, cash flows and financial condition.

The worldwide economic slowdown has and could continue to adversely affect our businesses and our profitability. If economic growth continues to remain slow, many customers may continue to delay purchases or reduce purchase quantities. This could result in the continued reduction in sales of certain of our products, slower adoption of both new technologies and upgrades to existing technologies and could also result in increased price competition. Continued market disruptions and broader economic downturns also increase our exposure to losses from bad debts. Among other affects we have seen during the slowdown, some of our customers, such as hospitals and healthcare systems in the United States, who rely on the credit or equity markets for access to capital, have and may continue to delay purchases of our products and services until the credit or equity markets recover. If economic or other factors cause financial institutions to fail, we could lose current or potential customers. During this period of uncertainty, we anticipate lower sales of patient monitoring, diagnostic cardiology and anesthesia systems products than we have historically experienced, resulting in a negative impact on our business, financial condition, results of operations, cash flows, strategies and prospects. We cannot predict when the world's financial markets will recover and therefore when this period of delayed and diminished purchasing will end. A prolonged delay could have a material adverse effect on our business, financial condition and results of operations. In addition, if the current turmoil in the financial markets continues, the variable interest rates payable under our credit facilities could be adversely affected or it could be more difficult to obtain or renew such facilities in the future. Any or all of these factors, as well as other consequences of the current economic conditions which cannot currently be anticipated, could have a material adverse effect on our revenues, earnings and cash flows and otherwise adversely affect our financial condition.

If we do not introduce new products in a timely manner, our products could become obsolete and our operating results would suffer.

We sell many of our products in industries characterized by rapid technological changes, frequent new product and service introductions and evolving industry standards and customer needs. Without the timely introduction of new products and enhancements, our products could become technologically obsolete over time, in which case our revenue and operating results would suffer. The success of our new product offerings will depend upon several factors, including our ability to:

- accurately anticipate customer needs;
- innovate and develop new technologies and applications;
- successfully commercialize new technologies in a timely manner;
- price our products competitively and manufacture and deliver our products in sufficient volumes and on time; and
- differentiate our offerings from our competitors' offerings.

Some of our products are used by our customers to develop, test and manufacture their products. We therefore must anticipate industry trends and develop products in advance of the commercialization of our customers' products. In developing any new product, we may be required to make a substantial investment

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before we can determine the commercial viability of the new product. If we fail to accurately foresee our customers' needs and future activities, we may invest heavily in research and development of products that do not lead to significant revenues.

Interruptions in our ability to purchase raw materials and subcomponents may adversely affect our profitability.

We purchase raw materials and certain subcomponents from third parties. Standard purchase order terms are as long as one year at fixed costs, but we do not have guaranteed long-term supply arrangements with our suppliers. In addition, for certain raw materials and subcomponents that we use there are a limited number of potential suppliers that we have qualified or that we are currently able to qualify. Consequently, some of the key raw materials and subcomponents that we use are currently available to us only from a single vendor. The reliance on a single qualified vendor could result delays in delivering products or increases in the cost of manufacturing the affected products. Any material interruption in our ability to purchase necessary raw materials or subcomponents could adversely affect our ability to fulfill customer orders and therefore could ultimately have a material adverse effect on our business, financial condition and results of operations.

We may not be able to successfully implement our acquisitions strategy, integrate acquired businesses into our existing business or make acquired businesses profitable.

One of our strategies is to supplement our internal growth by acquiring businesses and technologies that complement or augment our existing product lines. This growth has placed, and may continue to place, significant demands on our management, working capital and financial resources. We may be unable to identify or complete promising acquisitions for many reasons, including:

- competition among buyers;
- the need for regulatory approvals, including antitrust approvals; and
- the high valuations of businesses.

Some of the businesses we may seek to acquire may be marginally profitable or unprofitable. For these acquired businesses to achieve acceptable levels of profitability, we must improve their management, operations, products and market penetration. We may not be successful in this regard and may encounter other difficulties in integrating acquired businesses into our existing operations.

To finance our acquisitions, we may have to raise additional funds, through either public or private financings. We may be unable to obtain such funds or may be able to do so only on unfavorable terms.

Our acquisition and alliance activities could disrupt our ongoing business.

We intend to continue to make investments in companies, products and technologies, either through acquisitions, investments or alliances. Acquisition and alliance activities often involve risks, including:

- difficulty in assimilating the acquired operations and employees and realizing synergies expected to result from the acquisition;
- difficulty in managing product co-development activities with our alliance partners;
- difficulty in retaining the key employees of the acquired operation;
- disruption of our ongoing business;
- inability to successfully integrate the acquired technologies and operations into our businesses and maintain uniform standards, controls, policies and procedures; and
- lacking the experience necessary to enter into new product or technology markets successfully.

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Integrating acquired businesses has been and will continue to be complex, time consuming and expensive, and can negatively impact the effectiveness of our internal control over financial reporting. The use of debt to fund acquisitions or for other related purposes increases our interest expense and leverage. If we issue equity securities as consideration in an acquisition, current shareholders' percentage ownership and earnings per share may be diluted. As a result of these and other risks, we cannot be certain that our previous or future acquisitions will be successful and will not materially adversely affect the conduct, operating results or financial condition of our business.

Acquisition and alliance activities by our competitors could disrupt our ongoing business.

From time to time, our competitors acquire or enter into exclusive arrangements with companies with whom we do business or may do business in the future. Reductions in the number of partners with whom we may do business in a particular context may reduce our ability to enter into critical alliances on attractive terms or at all, and the termination of an existing alliance by a business partner may disrupt our operations.

Our ability to successfully adapt to ongoing organizational changes could impact our business results.

We have executed a number of significant business and organizational changes to rationalize our overall cost structure. These changes have included and may continue to include the implementation of cost-cutting measures and the consolidation of facilities. We expect these types of changes may continue from time to time in the future as we uncover additional opportunities to streamline our operations. Successfully managing these changes is critical to our productivity improvement and business success. If we are unable to successfully manage these changes, while continuing to invest in business growth, our financial results could be adversely impacted.

Economic, political and other risks associated with international sales and operations could adversely affect our sales.

In fiscal 2008, revenues from shipments made to customers outside of the United States accounted for approximately 47% of our revenues, 44% in fiscal 2009 and 43% in fiscal 2010. Of the revenues generated during fiscal 2010 from shipments made to customers outside of the United States, 29% represented sales made by subsidiaries based in the United States to foreign customers, and the balance represented sales generated by foreign subsidiaries. Since we sell certain of our products worldwide, our businesses are subject to risks associated with doing business internationally. We anticipate that revenues from international operations will continue to represent a substantial portion of our total revenue. In addition, many of our manufacturing facilities, and therefore employees, suppliers, real property, capital equipment, cash and other assets are located outside the United States. Accordingly, our future results could be harmed by a variety of factors, including:

- changes in foreign currency exchange rates;
- changes in a country's or region's political or economic conditions, particularly in developing or emerging markets;
- longer payment cycles of foreign customers and difficulty of collecting receivables in foreign jurisdictions;
- trade protection measures and import or export licensing requirements;
- differing legal and court systems;
- differing tax laws and changes in those laws;
- difficulty in staffing and managing widespread operations;
- difficulty in managing distributors and sales agents and their compliance with applicable laws;
- differing labor laws and changes in those laws;

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- differing protection of intellectual property and changes in that protection; and
- differing regulatory requirements and changes in those requirements.

Our competitors may seek to challenge the intellectual property rights on which our products are based.

As we introduce any new and potentially promising product, companies possessing competing technologies may be motivated to assert infringement claims in order to delay or diminish potential sales and challenge our right to market such product. In addition, we may find it necessary to initiate litigation in order to protect our patent or other intellectual property rights.

Under either circumstance, we may incur significant expenses. In addition, claims of third parties against us could result in awards of substantial damages or court orders that could effectively prevent us from making, using or selling our products in the United States or abroad.

Our ongoing success is dependent upon the continued availability of certain key employees.

We are dependent in our operations on the continued availability of the services of our employees, many of whom are individually key to our current and future success, and the availability of new employees to implement our growth plans. In particular, we are dependent upon the services of Deepak Chopra, our Chairman of the Board of Directors, President and Chief Executive Officer. The market for skilled employees is highly competitive, especially for employees in technical fields. While our compensation programs are intended to attract and retain the employees required for it to be successful, ultimately, we may not be able to retain the services of all of our key employees or a sufficient number to execute on our plans. In addition, we may not be able to continue to attract new employees as required.

Substantial government regulation in the United States and abroad may restrict our ability to sell our patient monitoring, diagnostic cardiology and anesthesia systems.

The FDA and comparable regulatory authorities in foreign countries extensively and rigorously regulate our patient monitoring, diagnostic cardiology and anesthesia systems, including related development activities and manufacturing processes. In the United States, the FDA regulates the introduction of medical devices as well as the manufacturing, labeling and record-keeping procedures for such products. We are required to:

- obtain clearance before we can market and sell medical devices;
- satisfy content requirements applicable to our labeling, sales and promotional materials;
- comply with manufacturing and reporting requirements; and
- undergo rigorous inspections.

Our future products may not obtain FDA clearance on a timely basis, or at all. Our patient monitoring, diagnostic cardiology and anesthesia systems must also comply with the laws and regulations of foreign countries in which we develop, manufacture and market such products. In general, the extent and complexity of medical device regulation is increasing worldwide. This trend is likely to continue and the cost and time required to obtain marketing clearance in any given country may increase as a result. Our products may not obtain any necessary foreign clearances on a timely basis, or at all.

Once any of our patient monitoring, diagnostic cardiology and anesthesia systems is cleared for sale, regulatory authorities may still limit the use of such product, prevent its sale or manufacture or require a recall or withdrawal of such product from the marketplace. Following initial clearance from regulatory authorities, we continue to be subject to extensive regulatory requirements. Government authorities can withdraw marketing

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clearance due to our failure to comply with regulatory standards or due to the occurrence of unforeseen problems following initial clearance. Ongoing regulatory requirements are wide-ranging and govern, among other things:

- annual inspections to retain a CE mark for sale of products in the European Union;
- product manufacturing;
- supplier substitution;
- product changes;
- process modifications;
- medical device reporting; and
- product sales and distribution.

Broad-based legislation addressing the affordability and availability medical services in the United States may reduce the amount of funding available to hospitals for purchases of patient monitoring, diagnostic cardiology and anesthesia systems.

The U.S. Congress has recently enacted new healthcare legislation, including the Children’s Health Insurance Reauthorization Act, the American Recovery and Reinvestment Act and the Affordable Care Act. The effects of this legislation are likely to be far-reaching as they require extensive revision to the current health insurance system in the United States. This legislation will significantly affect the ways in which doctors, hospitals, healthcare systems and health insurance companies are compensated for the services they provide. However, it is not clear at this time whether and to what extent these changes may impact the ability of hospitals and hospital networks to purchase the patient monitoring, diagnostic cardiology and anesthesia systems that we sell or if it will alter market-based incentives that hospitals and hospital networks currently face to continually improve, upgrade and expand their use of such equipment. While this legislation could adversely affect us, at this time we cannot predict the extent of any impact on our business or results of operations.

We are subject to various environmental regulations which may impose liability on us whether or not we knew of or caused the release of hazardous substances on or in our facilities.

We are subject to various foreign and U.S. federal, state and local environmental laws, ordinances and regulations relating to the use, storage, handling and disposal of certain hazardous substances and wastes used or generated in the manufacturing and assembly of our products. Under such laws, we may become liable for the costs of removal or remediation of certain hazardous substances or wastes that have been or are being disposed of offsite as wastes or that have been or are being released on or in our facilities. Such laws may impose liability without regard to whether we knew of, or caused, the release of such hazardous substances or wastes. Any failure by us to comply with present or future regulations could subject us to the imposition of substantial fines, suspension of production, alteration of manufacturing processes, or cessation of operations, any of which could have a material adverse effect on our business, financial condition and results of operations.

We may be exposed to potential risks relating to our internal controls over financial reporting and our ability to have our independent registered public accounting firm attest to these controls.

As directed by the Sarbanes-Oxley Act of 2002, the Securities and Exchange Commission adopted rules requiring public companies to include in their annual reports an assessment of the effectiveness of the company’s internal controls over financial reporting. In addition, the independent registered public accounting firm auditing a public company’s financial statements must attest to and report on the operating effectiveness of the company’s internal controls over financial reporting. We evaluate our internal controls over financial reporting in order to allow our management to report on, and our independent registered public accounting firm to attest to, our internal controls.

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We expect to continue to expend significant resources in complying with the documentation and testing procedures required by the Sarbanes-Oxley Act of 2002. However, there will remain an ongoing risk that we will not comply with all of its requirements.

If our independent registered public accounting firm differs from us in its interpretation of the requirements imposed on us by the Sarbanes-Oxley Act of 2002, or if it is not satisfied with our internal controls over financial reporting or with the level at which such controls are documented, operated or reviewed, we may be delayed in filing reports with the Securities and Exchange Commission, our independent registered public accounting firm may decline to attest to our management's assessment or it may issue an adverse report. In addition, if our independent registered public accounting firm is unable to rely on our internal controls over financial reporting in connection with its audit of our financial statements and if it is unable to devise alternative procedures in order to satisfy itself as to the material accuracy of our financial statements and related disclosures, it is possible that we could receive a qualified or adverse audit opinion in connection with those financial statements.

Accordingly, in the future we may not receive a favorable report from our independent registered public accounting firm regarding our internal controls over financial reporting and the operating effectiveness of our internal controls over financial reporting. If we identify material weaknesses in our internal controls over financial reporting that we cannot remediate in a timely manner or if we receive an adverse report from our independent registered public accounting firm with respect to our internal controls over financial reporting, investors and others may lose confidence in the reliability of our financial statements and the market for our Common Stock could be adversely affected.

A failure of a key information technology system, process or site could have a material adverse impact on our ability to conduct business.

We rely extensively on information technology systems to interact with our employees and our customers. These interactions include, but are not limited to, ordering and managing materials from suppliers, converting materials to finished products, shipping product to customers, processing transactions, summarizing and reporting results of operations, complying with regulatory, legal or tax requirements, and other processes necessary to manage our business. If our systems are damaged or cease to function properly due to any number of causes, ranging from the failures of third-party services provider, to catastrophic events to power outages to security breaches, and our business continuity plans do not effectively compensate on a timely basis, we may suffer interruptions in our ability to manage operations which may adversely impact our results of operations and/or financial condition.

We receive significant amounts of research and development funding for our security and inspection systems from government grants and contracts, but we may not continue to receive comparable levels of funding in the future.

The U.S. government currently plays an important role in funding the development of certain of our security and inspection systems and sponsoring their deployment at airports, ports, military installations and border crossings. However, in the future, additional research and development funds from the government may not be available to us. If the government fails to continue to sponsor our technologies we may have to expend more resources on product development or cease development of certain technologies, which could adversely affect our business. In addition, any future grants to our competitors may improve their ability to develop and market competing products and cause our customers to delay purchase decisions, which could harm our ability to market our products.

Our credit facility contains provisions that could restrict our ability to finance our future operations or engage in other business activities that may be in our interest.

Our credit facility contains a number of significant covenants that, among other things, limit our ability to:

- dispose of assets;
- incur certain additional indebtedness;

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- repay certain indebtedness;
- create liens on assets;
- pay dividends on our Common Stock;
- make certain investments, loans and advances;
- repurchase or redeem capital stock;
- make certain capital expenditures;
- engage in acquisitions, mergers or consolidations; and
- engage in certain transactions with subsidiaries and affiliates.

These covenants could limit our ability to plan for or react to market conditions, finance our operations, engage in strategic acquisitions or disposals or meet our capital needs or could otherwise restrict our activities or business plans. Our ability to comply with these covenants may be affected by events beyond our control. In addition, our credit facility also requires us to maintain compliance with certain financial ratios. Our inability to comply with the required financial ratios or covenants could result in an event of default under our credit facility. A default, if not cured or waived, may permit acceleration of our indebtedness. In addition, our lenders could terminate their commitments to make further extensions of credit under our credit facility. If our indebtedness is accelerated, we cannot be certain that we will have sufficient funds to pay the accelerated indebtedness or that we will have the ability to refinance accelerated indebtedness on terms favorable to us or at all.

Changes in our tax rates could affect our future financial results.

Our future effective tax rates could be favorably or unfavorably affected by changes in the valuation of our deferred tax assets and liabilities, or by changes in tax laws or their interpretation. In addition, we are subject to the continuous examination of our income tax returns by the Internal Revenue Service and other tax authorities. We regularly assess the likelihood of adverse outcomes resulting from these examinations to determine the adequacy of our provision for income taxes. There can be no assurance that the outcomes from these continuous examinations will not have an adverse effect on our operating results and financial condition.

In March 2010, significant reforms to the healthcare system were adopted as law in the United States. The law includes provisions that, among other things, reduce or limit Medicare reimbursement, require all individuals to have health insurance (with limited exceptions) and impose increased taxes. Specifically, the law requires the medical device industry to subsidize healthcare reform in the form of an excise tax on United States sales of most medical devices beginning in 2013. The excise tax will increase the Company's operating expenses. While the new law could reduce medical procedure volumes, lower reimbursement for the Company's products, and impact the demand for the Company's products or the prices at which the Company sells its products, at this time we cannot predict the extent of any impact on our business or results of operations.

Our Certificate of Incorporation and other agreements contain provisions that could discourage a takeover.

Our Certificate of Incorporation authorizes our Board of Directors to issue up to 10,000,000 shares of Preferred Stock in one or more series, to fix the rights, preferences, privileges and restrictions granted to or imposed upon any wholly unissued shares of Preferred Stock, to fix the number of shares constituting any such series and to fix the designation of any such series, without further vote or action by shareholders. The terms of any series of Preferred Stock, which may include economic rights senior to our Common Stock and special voting rights, could adversely affect the rights of the holders of our Common Stock and thereby reduce the value of our Common Stock. We have no present plans to issue shares of Preferred Stock. The issuance of Preferred Stock, coupled with the concentration of ownership in the directors and executive officers, could discourage certain types of transactions involving an actual or potential change in control of our company, including

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transactions in which the holders of Common Stock might otherwise receive a premium for their shares over then current prices, otherwise dilute the rights of holders of Common Stock and may limit the ability of such shareholders to cause or approve transactions which they may deem to be in their best interests, all of which could have a material adverse effect on the market price of our Common Stock.

Our Certificate of Incorporation limit the liability of our directors, which may limit the remedies we or our shareholders have available.

Our Certificate of Incorporation provides that, pursuant to the Delaware General Corporation Law, the liability of our directors for monetary damages shall be eliminated to the fullest extent permissible under Delaware law, as that law exists currently and as it may be amended in the future. This is intended to eliminate the personal liability of a director for monetary damages in an action brought by us, or in our right for breach of a director's duties to us or our shareholders and may limit the remedies available to us or our shareholders. Under Delaware law, this provision does not apply to eliminate or limit a director's monetary liabilities for: (i) breaches of the director's duty of loyalty to us or our shareholders; (ii) acts or omissions not in good faith or which involve intentional misconduct or knowing violations of law; (iii) the unlawful payment of dividends or unlawful stock repurchases or redemptions under Section 174 of the Delaware General Corporation Law or (iv) transactions in which the director received an improper personal benefit. Additionally, under Delaware law, this provision does not limit a director's liability for the violation of, or otherwise relieve us or our directors from complying with, federal or state securities laws, nor does it limit the availability of non-monetary remedies such as injunctive relief or rescission for a violation of federal or state securities laws.

ITEM 1B. UNRESOLVED STAFF COMMENTS

None.

ITEM 2. PROPERTIES

As of June 30, 2010, we owned four facilities. The following table lists these facilities:

<u>Location</u>	<u>Description of Facility</u>	<u>Approximate Square Footage</u>
Hawthorne, California	Corporate headquarters and administrative, manufacturing, engineering, sales and marketing and service for our Optoelectronics and Manufacturing division	88,000
Surrey, England	Manufacturing, engineering, sales and marketing and service for our Security and Healthcare divisions	59,000
Batam, Indonesia	Manufacturing for our Optoelectronics and Manufacturing division	59,000
Ocean Springs, Mississippi	Manufacturing, engineering, sales and marketing and service for our Security and Optoelectronics and Manufacturing divisions	19,000

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As of June 30, 2010, we leased all of our other facilities. The following table lists the principal (*i.e.*, facilities greater than 50,000 square feet) physical properties that we lease:

<u>Location</u>	<u>Description of Facility</u>	<u>Approximate Square Footage</u>	<u>Expiration</u>
Camarillo, California	Manufacturing, engineering, sales and marketing and service for our Optoelectronics and Manufacturing division	60,000	2015
Sunnyvale, California	Manufacturing, engineering, sales and marketing and service for our Security division	62,500	2012
Torrance, California	Manufacturing, engineering, sales and marketing and service for our Security division	91,900	2012
Apex, North Carolina	Manufacturing, engineering, sales and marketing and service for our Security division	122,400	2012
Issaquah, Washington (1)	Manufacturing, engineering, sales and marketing and service for our Healthcare division	202,600	2014
Suzhou, China	Manufacturing, engineering, sales and marketing and service for our Healthcare division	53,000	2012
Hyderabad, India (2)	Manufacturing and engineering for our Security, Healthcare and Optoelectronics and Manufacturing divisions	55,300	2013
Johor Bahru, Malaysia	Manufacturing, engineering sales and service for our Security division	87,100	2012
Johor Bahru, Malaysia (3)	Manufacturing, engineering sales and service for our Optoelectronics and Manufacturing division	76,000	2010
Stoke on Trent, United Kingdom	Manufacturing, engineering, sales and marketing and service for our Security division	65,000	2020

- (1) The lease of the 202,600 square foot facility in Issaquah, Washington is composed of two leases in the same facility. One lease covers a 107,000 square foot area within the facility and the other covers a 95,600 square foot area within the facility. Both leases expire in December 2014.
- (2) The lease of the 55,300 square foot facility in Hyderabad, India comprises seven leases, ranging in size between 1,100 square feet and 19,800 square feet. Each of these leases expires in 2013.
- (3) Upon expiration of lease of the 76,000 facility in Johor Bahru, Malaysia we expect to renew the lease on the same or similar terms or, in the alternative, to move into another nearby facility.

We believe that our facilities are in good condition and are adequate to support our operations for the foreseeable future. We currently anticipate that we will be able to renew the leases that are scheduled to expire in the next few years on terms that are substantially the same as or better than those currently in effect. However,

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even if we were not able to renew one or more of the leases, we believe that suitable substitute space is available to relocate any of the facilities. Accordingly, we do not believe that our failure to renew any of the leases that are scheduled to expire in the next few years will have a material adverse effect on our operations.

ITEM 3. LEGAL PROCEEDINGS

We are involved in various claims and legal proceedings arising out of the ordinary course of business. In our opinion after consultation with legal counsel, the ultimate disposition of such proceedings will not likely have a material adverse effect on our financial position, future results of operations, or cash flows. In accordance with accounting standards related to contingencies, we have not accrued for loss contingencies relating to such matters because we believe that, although unfavorable outcomes in the proceedings may be possible, they are not considered by management to be probable or reasonably estimable. If one or more of these matters are resolved in a manner adverse to us, the impact on our results of operations, financial position and/or liquidity could be material.

ITEM 4. RESERVED

PART II**ITEM 5. MARKET FOR REGISTRANT'S COMMON EQUITY, RELATED STOCKHOLDER MATTERS AND ISSUER PURCHASES OF EQUITY SECURITIES****Stock Market and Other Information**

Our Common Stock is traded on The NASDAQ Global Market under the symbol "OSIS."

The following table sets forth the high and low sale prices of a share of our Common Stock as reported by The NASDAQ Global Market on a quarterly basis for fiscal 2009 and 2010. The prices shown reflect inter-dealer prices, without retail markup, markdown or commission and may not necessarily represent actual transactions.

<u>2009:</u>	<u>High</u>	<u>Low</u>
Quarter ended September 30, 2008	\$25.56	\$19.95
Quarter ended December 31, 2008	\$23.57	\$10.15
Quarter ended March 31, 2009	\$18.38	\$11.01
Quarter ended June 30, 2009	\$21.13	\$14.93
<u>2010:</u>	<u>High</u>	<u>Low</u>
Quarter ended September 30, 2009	\$21.23	\$16.31
Quarter ended December 31, 2009	\$28.49	\$17.03
Quarter ended March 31, 2010	\$32.58	\$25.21
Quarter ended June 30, 2010	\$30.43	\$22.80

As of August 23, 2010, there were approximately 75 holders of record of our Common Stock. This number does not include beneficial owners holding shares through nominees or in "street" name.

Dividend Policy

We have not paid any cash dividends since the consummation of our initial public offering in 1997 and anticipate that we will retain any available funds for use in the operation of our business. We do not currently intend to pay any cash dividends in the foreseeable future. Our Board of Directors will determine the payment of future cash dividends, if any. Certain of our current bank credit facilities restrict the payment of cash dividends and future borrowings may contain similar restrictions.

Issuer Purchases of Equity Securities

In March 1999, our Board of Directors authorized a stock repurchase program for the repurchase of up to 2 million shares of our Common Stock. In September 2004, we increased the number of shares available for repurchase under the stock repurchase program by 1 million shares. During the 12 months ended June 30, 2009, we repurchased 619,768 shares under this program. During the twelve months ended June 30, 2010, we did not repurchase any shares under this program. As of June 30, 2010, 711,205 shares were available for additional repurchase under the program. Upon repurchase, the shares were restored to the status of authorized but unissued and we recorded them as a reduction in the number of shares of Common Stock issued and outstanding in our Consolidated Financial Statements.

[Table of Contents](#)**Equity Compensation Plans**

The following table provides information concerning our equity compensation plans as of June 30, 2010.

<u>Plan category</u>	<u>Number of securities to be issued upon exercise of outstanding options, warrants and rights</u> (a)	<u>Weighted-average exercise price of outstanding options, warrants and rights</u> (b)	<u>Number of securities remaining available for future issuance under equity compensation plans (excluding securities reflected in column (a))</u> (c)
Equity compensation plans approved by security holders (1)(2)	1,556,110	17.46	524,017
Equity participation plans not approved by security holders	—	N/A	—
Total	<u>1,556,110</u>	<u>17.46</u>	<u>524,017</u>

(1) Includes shares of our Common Stock issuable upon exercise of options under our 2006 Equity Participation Plan.

(2) Of the 524,017 securities remaining available for future issuance under our 2006 Equity Participation Plan, 349,121 shares are available to be issued as restricted stock.

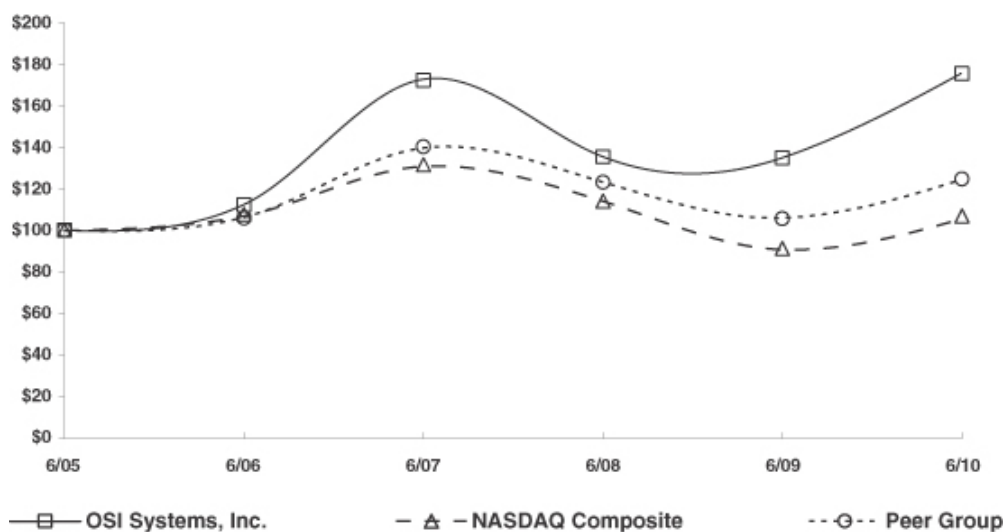
Performance Graph

The graph below compares the cumulative total shareholder return for the period beginning on the market close on the last trading day before the beginning of our fifth preceding fiscal year through and including the end of our last completed fiscal year, with (a) The NASDAQ Global Market Index and (b) a peer group of publicly-traded issuers with which we have generally competed.

The peer group includes the following companies: American Science & Engineering (AMEX Symbol: ASE) and Analogic Corporation (NASDAQ Symbol: ALOG).

The graph assumes that \$100.00 was invested on June 30, 2005 in (a) our Common Stock, (b) The NASDAQ Global Market Index and (c) the companies comprising the peer group described above (weighted according to each respective issuer's stock market capitalization at the beginning of each period for which a return is indicated). The graph assumes that all dividends were reinvested. Historical stock price performance is not necessarily indicative of future stock price performance.

Comparison of 5 Year Cumulative Total Return*
Assumes Initial Investment of \$100
June 2005 through June 2010
Among OSI Systems, Inc.,
The NASDAQ Composite Index And A Peer Group



* \$100 invested on June 30, 2005 in stock & index-including reinvestment of dividends.

The following table provides the same information in tabular form as of June 30:

	2005	2006	2007	2008	2009	2010
OSI Systems, Inc.	100.00	112.54	173.21	135.66	135.07	175.87
The NASDAQ Composite Index	100.00	107.09	131.06	114.12	90.80	105.51
Peer Group	100.00	106.36	140.00	123.53	106.12	124.48

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ITEM 6. SELECTED FINANCIAL DATA

The following table sets forth our selected consolidated financial data as of and for each of the five fiscal years ended June 30, 2010, and is derived from our Consolidated Financial Statements. The Consolidated Financial Statements as of June 30, 2009 and 2010, and for each of the years in the three-year period ended June 30, 2010, are included elsewhere in this report. The following data should be read in conjunction with “Management’s Discussion and Analysis of Financial Condition and Results of Operations” and the Consolidated Financial Statements and Notes thereto included elsewhere in this report.

	Year Ended June 30,				
	2006	2007	2008	2009	2010
	(in thousands, except earnings per share data)				
Consolidated Statements of Operations Data (1):					
Revenues	\$ 452,686	\$ 532,284	\$ 623,088	\$ 590,361	\$ 595,111
Cost of goods sold	276,025	354,067	404,049	388,910	377,077
Gross profit	176,661	178,217	219,039	201,451	218,034
Operating expenses:					
Selling, general and administrative	140,823	151,031	150,082	137,985	139,830
Research and development	35,839	44,446	45,361	36,862	38,577
Restructuring and other charges	800	26,071	4,688	7,123	2,859
Total operating expenses	177,462	221,548	200,131	181,970	181,266
Income (loss) from operations	(801)	(43,331)	18,908	19,481	36,768
Interest expense and other income, net	467	(11,697)	4,469	2,936	1,772
Income (loss) before income taxes	(1,268)	(31,634)	14,439	16,545	34,996
Provision (benefit) for income taxes	1,090	(12,876)	579	5,393	11,439
Net income (loss)	\$ (2,358)	\$ (18,758)	\$ 13,860	\$ 11,152	\$ 23,557
Net income (loss) available to common shareholders—diluted	\$ (2,738)	\$ (18,815)	\$ 13,860	\$ 11,152	\$ 23,557
Basic earnings (loss) per common share	\$ (0.14)	\$ (1.11)	\$ 0.80	\$ 0.64	\$ 1.32
Diluted earnings (loss) per common share	\$ (0.17)	\$ (1.12)	\$ 0.78	\$ 0.63	\$ 1.28
Weighted average shares outstanding—diluted	16,517	16,844	17,735	17,596	18,389

	Year Ended June 30,				
	2006	2007	2008	2009	2010
	(in thousands)				
Consolidated Balance Sheet Data (1):					
Cash and cash equivalents	\$ 13,799	\$ 15,980	\$ 18,232	\$ 25,172	\$ 51,989
Working capital	162,156	158,741	194,958	187,608	204,607
Total assets	403,498	451,483	507,641	474,828	513,114
Long-term debt	5,483	25,709	49,091	39,803	23,366
Total debt	17,591	48,228	74,341	52,360	36,109
Total shareholders’ equity	248,947	247,212	278,021	276,000	313,710

(1) Results of operations for fiscal years 2006 through 2010, and our financial position as of June 30, 2006, 2007, 2008, 2009 and 2010 incorporate the effect of several acquisitions.

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

Overview

We are a vertically integrated designer and manufacturer of specialized electronic systems and components for critical applications. We sell our products in diversified markets, including homeland security, healthcare, defense and aerospace. We have three operating divisions: (a) Security, providing security and inspection systems; (b) Healthcare, providing patient monitoring, diagnostic cardiology and anesthesia systems; and (c) Optoelectronics and Manufacturing, providing specialized electronic components for affiliated end-products divisions, as well as for applications in the defense and aerospace markets, among others.

Security Division. Through our Security division, we design, manufacture and market security and inspection systems worldwide for sale primarily to U.S. and foreign government agencies, and provide turnkey security screening solutions. These products and services are used to inspect baggage, cargo, vehicles and other objects for weapons, explosives, drugs and other contraband as well as to screen people. Revenues from our Security division accounted for 42% of our total consolidated revenues for fiscal 2010.

Healthcare Division. Through our Healthcare division, we design, manufacture and market patient monitoring, diagnostic cardiology and anesthesia systems worldwide for sale primarily to hospitals and medical centers. Our products monitor patients in critical, emergency and perioperative care areas of the hospital and provide such information, through wired and wireless networks, to physicians and nurses who may be at the patient's bedside, in another area of the hospital or even outside the hospital. Revenues from our Healthcare division accounted for 35% of our total consolidated revenues for fiscal 2010.

In October 2005, Spacelabs Healthcare, Inc., a subsidiary comprising the business operations of our Healthcare division, completed an initial public offering of approximately 20% of its total issued and outstanding common stock. The Spacelabs Healthcare shares traded under the ticker symbol "SLAB" on the AIM (formerly known as the Alternative Investment Market), a stock market administered by the London Stock Exchange. In the second quarter of fiscal 2007, we began repurchasing publicly-traded shares of Spacelabs Healthcare, increasing our ownership to 84% as of June 30, 2007. By December 31, 2007, we increased our ownership in Spacelabs Healthcare to 100% by repurchasing all remaining shares of Spacelabs Healthcare. Effective January 24, 2008, we cancelled Spacelabs Healthcare's AIM listing.

Optoelectronics and Manufacturing Division. Through our Optoelectronics and Manufacturing division, we design, manufacture and market optoelectronic devices and provide electronics manufacturing services worldwide for use in a broad range of applications, including aerospace and defense electronics, security and inspection systems, medical imaging and diagnostics, computed tomography (CT), telecommunications, office automation, computer peripherals and industrial automation. We also provide our optoelectronic devices and value-added manufacturing services to our own Security and Healthcare divisions. Revenues from our Optoelectronics and Manufacturing division accounted for approximately 23% of our total consolidated revenues for fiscal 2010.

Consolidated Results.

Fiscal 2010 Compared with Fiscal 2009. We reported consolidated operating profit of \$36.8 million for fiscal 2010, an 89% improvement from the \$19.5 million operating profit reported for fiscal 2009. This improved profitability was driven primarily by a \$16.5 million improvement in gross profit as a result of product mix, operational efficiencies gained in our supply chain and ongoing restructuring and cost-cutting initiatives that we have undertaken during the past three fiscal years as well as a reduction in non-recurring restructuring and other charges of \$4.3 million to \$2.9 million in fiscal 2010 from the \$7.1 million incurred in fiscal 2009. These improvements in operating profitability were partially offset by a \$3.6 million increase in selling, general and administrative (SG&A) and research and development (R&D) expenses.

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Fiscal 2009 Compared with Fiscal 2008. We reported consolidated operating profit of \$19.5 million for fiscal 2009, a 3% improvement from the \$18.9 million operating profit reported for fiscal 2008. We realized this \$0.6 million increase in operating profits despite a 5% decrease in total revenue in fiscal 2009 to \$590.4 million from \$623.1 million in fiscal 2008. In addition, we incurred \$7.1 million of non-recurring restructuring and other charges in fiscal 2009, an increase of \$2.4 million from the \$4.7 million incurred in fiscal 2008. This improved profitability was driven primarily by a \$20.6 million reduction in SG&A and R&D expenses following restructuring and cost-cutting initiatives we undertook during fiscal 2008 and 2009. This reduction in operating expenses more than offset a \$17.5 million reduction in gross profit, which resulted from lower revenues, primarily in our Healthcare division, which saw revenues decrease by 17%.

Acquisitions. Historically, an active acquisition program has been an important element of our corporate strategy. However, in recent years, we slowed our acquisition activity to conduct a global review of our core businesses and technologies and to fully integrate the acquisitions we made during the past several years. This global review and integration resulted in significant restructuring and other charges in fiscal 2008 through 2010. See Note 7 to Consolidated Financial Statements for further discussion. Looking forward, we continue to believe that an active acquisition program supports our long-term strategic direction. We look to acquisitions to strengthen our competitive position, expand our customer base and augment our considerable research and development programs. Through such efforts we aim to accelerate innovation, improve earnings and increase overall stockholder value.

Critical Accounting Policies and Estimates

Our discussion and analysis of our financial condition and results of operations is based on our Consolidated Financial Statements, which have been prepared in conformity with accounting principles generally accepted in the United States. Our preparation of these Consolidated Financial Statements requires us to make judgments and estimates that affect the reported amounts of assets and liabilities, disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period. We base our estimates on historical experience and on various other assumptions that we believe to be reasonable under the circumstances. As a result, actual results may differ from such estimates. Our senior management has reviewed these critical accounting policies and related disclosures with the Audit Committee of our Board of Directors. The following summarizes our critical accounting policies and significant estimates used in preparing our Consolidated Financial Statements:

Revenue Recognition. We recognize revenue upon shipment of products when title and risk of loss passes, and when terms are fixed and collection is probable. In cases where installation services are essential to the functionality of the equipment, we defer the portion of revenue for the sale attributable to installation until we have completed the installation. When terms of sale include subjective customer acceptance criteria, we defer revenue until the acceptance criteria are met. Concurrent with the shipment of the product, we accrue estimated product return reserves and warranty expenses. Critical judgments made by management related to revenue recognition include the determination of whether or not customer acceptance criteria are perfunctory or inconsequential. The determination of whether or not the customer acceptance terms are perfunctory or inconsequential impacts the amount and timing of the revenue that we recognize. Critical judgments also include estimates of warranty reserves, which are established based on historical experience and knowledge of the product.

We recognize revenues from separate service maintenance contracts ratably over the term of the agreements. For other services, we recognize service revenues as we perform the services. Deferred revenue for services arises from advance payments received from customers for services not yet performed. We record billed shipping and handling fees as revenue and the associated costs as cost of goods sold.

Allowance for Doubtful Accounts. The allowance for doubtful accounts involves estimates based on management's judgment, review of individual receivables and analysis of historical bad debts. We monitor

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collections and payments from our customers and we maintain allowances for doubtful accounts for estimated losses resulting from the inability of our customers to make required payments. We also assess current economic trends that might impact the level of credit losses in the future. If the financial condition of our customers were to deteriorate, resulting in an impairment of their ability to make payments, additional allowances could be required.

Inventory. Inventory is stated at the lower of cost or market. Cost is determined on the first-in, first-out method. We write down inventory for slow-moving and obsolete inventory based on assessments of future demands, market conditions and customers who may be experiencing financial difficulties. If these factors were to become less favorable than those projected, additional inventory write-downs could be required.

Income Taxes. Our annual tax rate is based on our income, statutory tax rates and tax planning opportunities available to us in the various jurisdictions in which we operate. Tax laws are complex and subject to different interpretations by the taxpayer and respective governmental taxing authorities. Significant judgment is required in determining our tax expense and in evaluating our tax positions including evaluating uncertainties. We review our tax positions quarterly and adjust the balances as new information becomes available.

Deferred income tax assets represent amounts available to reduce income taxes payable on taxable income in future years. Such assets arise because of temporary differences between the financial reporting and tax bases of assets and liabilities, as well as from net operating loss and tax credit carryforwards. We evaluate the recoverability of these future tax deductions by assessing the adequacy of future expected taxable income from all sources, including reversal of taxable temporary differences, forecasted operating earnings and available tax planning strategies. These sources of income inherently rely on estimates. To provide insight, we use our historical experience and our short and long-range business forecasts. We believe it is more likely than not that a portion of the deferred income tax assets may expire unused and have established a valuation allowance against them. Although realization is not assured for the remaining deferred income tax assets, we believe it is more likely than not that the deferred tax assets will be fully recoverable within the applicable statutory expiration periods. However, deferred tax assets could be reduced in the near term if our estimates of taxable income are significantly reduced or available tax planning strategies are no longer viable.

Business Combinations. Under the purchase method of accounting, we allocate the purchase price of acquired companies to the tangible and identifiable intangible assets acquired and liabilities assumed based on their estimated fair values. We record the excess of purchase price over the aggregate fair values as goodwill. We engage third-party appraisal firms to assist us in determining the fair values of assets acquired and liabilities assumed. These valuations require us to make significant estimates and assumptions, especially with respect to intangible assets. Critical estimates in valuing purchased technology, customer lists and other identifiable intangible assets include future cash flows that we expect to generate from the acquired assets. If the subsequent actual results and updated projections of the underlying business activity change compared with the assumptions and projections used to develop these values, we could experience impairment charges. In addition, we have estimated the economic lives of certain acquired assets and these lives are used to calculate depreciation and amortization expense. If our estimates of the economic lives change, depreciation or amortization expenses could be accelerated or slowed.

Impairment of Long-Lived Assets. We test goodwill for impairment at the reporting unit level at least annually and more frequently upon the occurrence of certain events. For purposes of testing for goodwill impairment, we have determined that we have three reporting units for goodwill impairment review purposes, consisting of our Security division, our Healthcare division and our Optoelectronics and Manufacturing division. We test goodwill for impairment annually during the second fiscal quarter using a two-step process. First, we determine if the carrying amount of any of the reporting units exceeds its fair value. The fair value of the reporting units was calculated using the income approach and the market approach. Under the income approach, the fair value of the reporting units was calculated by estimating the present value of associated future cash flows. Under the market approach, the fair value was calculated using the guideline public company method and

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the mergers and acquisitions method. We performed this annual impairment test for goodwill during the second quarter of fiscal 2010 and concluded that there was no impairment of goodwill.

We evaluate long-lived assets, including intangible assets other than goodwill, for impairment whenever events or changes in circumstances indicate that the carrying value of an asset may not be recoverable. An impairment is considered to exist if the total estimated future cash flows on an undiscounted basis are less than the carrying amount of the assets. If an impairment does exist, we measure the impairment loss and record it based on discounted estimated future cash flows. In estimating future cash flows, we group assets at the lowest level for which there are identifiable cash flows that are largely independent of cash flows from other asset groups. Our estimate of future cash flows is based upon, among other things, certain assumptions about expected future operating performance, growth rates and other factors.

Although we believe the assumptions and estimates we have made in the past have been reasonable and appropriate, different assumptions and estimates could materially impact our reported financial results. More conservative estimates of the anticipated future benefits from these businesses could result in impairment charges, which would decrease net income and result in lower asset values on our balance sheet.

Stock-Based Compensation Expense. We account for share based compensation using fair value recognition provisions. Thus, we record stock-based compensation as a charge to earnings net of the estimated impact of forfeited awards. As such, we recognize stock-based compensation cost only for those stock-based awards that are estimated to ultimately vest over their requisite service period, based on the vesting provisions of the individual grants.

The process of estimating the fair value of stock-based compensation awards and recognizing stock-based compensation cost over their requisite service period involves significant assumptions and judgments. We estimate the fair value of stock option awards on the date of grant using the Black-Scholes option-valuation model which requires that we make certain assumptions regarding: (i) the expected volatility in the market price of our Common Stock; (ii) dividend yield; (iii) risk-free interest rates; and (iv) the period of time employees are expected to hold the award prior to exercise (referred to as the expected holding period). We estimate the fair value of restricted stock awards on the date of the grant using the market price of our Common Stock on that date. In addition, we are required to estimate the expected impact of forfeited awards and recognize stock-based compensation cost only for those awards expected to vest. If actual forfeiture rates differ materially from our estimates, stock-based compensation expense could differ significantly from the amounts we have recorded in the current period. We periodically review actual forfeiture experience and revise our estimates, as necessary. We recognize the cumulative effect on current and prior periods of a change in the estimated forfeiture rate as compensation cost in earnings in the period of the revision. As a result, if we revise our assumptions and estimates, our stock-based compensation expense could change materially in the future. See Note 9 (Stock-based Compensation) to the Consolidated Financial Statements for a further discussion of stock-based compensation.

Legal and Other Contingencies. We are subject to various claims and legal proceedings. Each fiscal quarter, we review the status of each significant legal dispute to which we are a party and assess our potential financial exposure, if any. If the potential financial exposure from any claim or legal proceeding is considered probable and the amount can be reasonably estimated, we record a liability and an expense for the estimated loss. Significant judgment is required in both the determination of probability and the determination as to whether an exposure is reasonably estimable. Because of uncertainties related to these matters, accruals are based only on the best information available at the time. As additional information becomes available, we reassess the potential liability related to our pending claims and litigation and revise our estimates accordingly. Such revisions in the estimates of the potential liabilities could have a material impact on our results of operations and financial position.

[Table of Contents](#)**Net Revenues**

The table below and the discussion that follows are based upon the way we analyze our business. See Note 15 to the Consolidated Financial Statements for additional information about business segments.

	<u>2008</u>	<u>% of Net Sales</u>	<u>2009</u>	<u>% of Net Sales</u>	<u>2010</u>	<u>% of Net Sales</u>	<u>2008-2009 % Change</u>	<u>2009-2010 % Change</u>
					(Dollars in millions)			
Security	\$225.8	36%	240.9	41%	251.5	42%	7%	4%
Healthcare	256.7	41%	214.3	36%	206.6	35%	(17)%	(4)%
Optoelectronics / Manufacturing	140.6	23%	135.2	23%	137.0	23%	(4)%	1%
Total Sales	<u>\$623.1</u>		<u>\$590.4</u>		<u>\$595.1</u>		<u>(5)%</u>	<u>1%</u>

Fiscal 2010 Compared with Fiscal 2009. Net revenues for fiscal 2010 increased \$4.7 million, or 1%, to \$595.1 million from \$590.4 million for fiscal 2009.

Revenues for the Security division for fiscal 2010 increased \$10.6 million, or 4%, to \$251.5 million, from \$240.9 million for fiscal 2009. The increase was attributable to a \$6.1 million, or 3%, increase in equipment sales, driven by a \$25.5 million increase in people screening and hold baggage screening products, which was partially offset by a \$19.4 million decrease in revenues from our cargo inspection products. In addition, service revenues increased by \$4.5 million or 10%, due to the growing installed base of products from which we derive service revenue as warranty periods expire.

Revenues for the Healthcare division for fiscal 2010 decreased \$7.7 million, or 4%, to \$206.6 million, from \$214.3 million for fiscal 2009. The decrease was primarily attributable to an 18% decrease in our anesthesia equipment sales and a \$6.8 million reduction in the revenues of other product lines such as pulse oximeters. We believe that these decreases were primarily due to generally poor overall economic conditions particularly in Europe. This decrease was partially offset by a \$2.8 million increase in patient monitoring revenues mainly in North America.

Revenues for the Optoelectronics and Manufacturing division for fiscal 2010 increased \$1.8 million, or 1.4%, to \$137.0 million from \$135.2 million for fiscal 2009. This increase was driven by an increase in our contract manufacturing business of \$4.7 million, and partially offset by lower commercial optoelectronics sales, which decreased by \$2.8 million.

Fiscal 2009 Compared with Fiscal 2008. Net revenues for fiscal 2009 decreased \$32.7 million, or 5%, to \$590.4 million from \$623.1 million for fiscal 2008. Excluding the adverse impact of foreign exchange, the decrease in revenues from fiscal 2008 to 2009 would have been approximately 3%.

Revenues for the Security division for fiscal 2009 increased \$15.1 million, or 7%, to \$240.9 million, from \$225.8 million for fiscal 2008. The increase was attributable to a 5% increase in sales of baggage and parcel inspection, people screening and hold baggage screening equipment; a 3% increase in sales of cargo and vehicle inspection systems equipment; and a 20% increase in service revenue. The Security division achieved this growth despite the impact of a worldwide economic slowdown. We believe that such increases in revenues reflect continuing worldwide acceptance of our Security division's products.

Revenues for the Healthcare division for fiscal 2009 decreased \$42.4 million, or 17%, to \$214.3 million, from \$256.7 million for fiscal 2008. The decrease was primarily attributable to: (i) a 16%, decrease in patient monitoring revenues, primarily in North America; (ii) a reduction in diagnostic cardiology equipment sales of 21% and (iii) lower clinical trials services sales of 55%. We believe that such decreases were a consequence of the severe downturn in world financial markets that occurred during fiscal 2009 and the inability of some of our customers, who rely on the credit or equity markets for access to capital, to fund purchases of our products and services.

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Revenues for the Optoelectronics and Manufacturing division for fiscal 2009 decreased \$5.4 million, or 4%, to \$135.2 million, from \$140.6 million for fiscal 2008. The decrease was primarily attributable to lower weapons simulation sales of \$4.6 million as well as lower commercial optoelectronics sales of \$7.7 million, partially offset by an increase in contract manufacturing sales of \$5.9 million, including product shipments under a significant defense-industry related contract.

Gross Profit

	<u>2008</u>	<u>% of Net Sales</u>	<u>2009</u>	<u>% of Net Sales</u>	<u>2010</u>	<u>% of Net Sales</u>
	(Dollars in millions)					
Gross profit	\$219.0	35.1%	\$201.5	34.1%	\$218.0	36.6%

Fiscal 2010 Compared with Fiscal 2009. Gross profit increased \$16.5 million, or 8%, to \$218.0 million for fiscal 2010, from \$201.5 million for fiscal 2009. The gross margin increased to 36.6% in fiscal 2010 from 34.1% in fiscal 2009. This increase was primarily attributable to manufacturing efficiencies gained through facility consolidation and operational improvement initiatives and due to a favorable product mix within our Security and Healthcare divisions.

Fiscal 2009 Compared with Fiscal 2008. Gross profit decreased \$17.5 million, or 8%, to \$201.5 million for fiscal 2009, from \$219.0 million for fiscal 2008. The gross margin decreased to 34.1% in fiscal 2009 from 35.1% in fiscal 2008. The decrease in gross profit is primarily a result of a 5% decrease in sales and changes in our product mix, most notably the 17% decrease in revenues in our Healthcare division (products sold by our Healthcare division generally carry higher gross margins than products sold by our other divisions) and an increase in contract manufacturing sales by our Optoelectronics and Manufacturing division, which sales generally carry lower gross margins than most of our other product offerings. The negative impact on our gross margins by this product mix was partially offset by manufacturing efficiencies gained through facility consolidations and operational improvement initiatives undertaken during the past several years.

Operating Expenses

	<u>2008</u>	<u>% of Net Sales</u>	<u>2009</u>	<u>% of Net Sales</u>	<u>2010</u>	<u>% of Net Sales</u>	<u>2008-2009 % Change</u>	<u>2009-2010 % Change</u>
	(Dollars in millions)							
Selling, general and administrative	\$150.1	24.1%	\$137.9	23.4%	\$139.8	23.5%	(8)%	1%
Research and development	45.3	7.3%	36.9	6.2%	38.6	6.5%	(19)%	5%
Restructuring and other charges	4.7	0.7%	7.1	1.2%	2.9	0.5%	53%	(61)%
Total operating expenses	<u>\$200.1</u>	<u>32.1%</u>	<u>\$181.9</u>	<u>30.8%</u>	<u>\$181.3</u>	<u>30.5%</u>	<u>(9)%</u>	<u>— %</u>

Selling, General and Administrative

Fiscal 2010 Compared with Fiscal 2009. Selling, general and administrative expenses consisted primarily of compensation paid to sales, marketing and administrative personnel, professional service fees and marketing expenses. For fiscal 2010, SG&A expenses increased by \$1.9 million, or 1%, to \$139.8 million, from \$137.9 million for fiscal 2009. This increase in spending was primarily attributable to an increase in SG&A costs to support revenue growth in the Security division, partially offset by lower spending throughout other parts of the company, especially in the Healthcare division, as a result of cost containment initiatives we have been undertaking to better leverage our cost structure.

Fiscal 2009 Compared with Fiscal 2008. For fiscal 2009, SG&A expenses decreased by \$12.2 million, or 8%, to \$137.9 million, from \$150.1 million for fiscal 2008. This reduction in spending was a direct result of our cost containment initiatives and restructuring activities throughout the Company, which were heavily focused in our Healthcare division. Due to these increased efficiencies, our SG&A as a percentage of sales decreased to 23.4% in fiscal 2009 as compared to 24.1% in fiscal 2008, despite the 5% sales decrease.

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Research and Development

Our Security and Healthcare divisions have historically invested substantial amounts in research and development. We intend to continue this trend in future years, although specific programs may or may not continue to be funded and funding levels may fluctuate.

Fiscal 2010 Compared with Fiscal 2009. Research and development expenses included research related to new product development and product enhancement expenditures. For fiscal 2010, such expenses increased by \$1.7 million, or 5%, to \$38.6 million, from \$36.9 million for fiscal 2009. As a percentage of revenues, R&D expenses were 6.5% in fiscal 2010, compared to 6.2% in fiscal 2009. The increase in R&D spending in fiscal 2010 resulted from an increase in both our Security and Healthcare divisions in support of new product introductions.

Fiscal 2009 Compared with Fiscal 2008. For fiscal 2009, R&D expenses decreased by \$8.4 million, or 19%, to \$36.9 million, from \$45.3 million for fiscal 2008. As a percentage of revenues, R&D expenses were 6.2% in fiscal 2009, compared to 7.3% in fiscal 2008. The decrease in R&D spending was primarily attributable to cost reduction efforts in our Healthcare division and increased levels of government R&D funding to our Security division.

Restructuring and Other Charges

Beginning in fiscal 2007 and continuing through fiscal 2009, we initiated a series of restructuring activities that were intended to realign our global capacity and infrastructure with demand by our customers and fully integrate acquisitions made in the last several years, thereby improving our operational efficiency. These activities included reducing excess workforce and capacity, consolidating and relocating certain manufacturing facilities and reviewing the value of certain technologies and product lines. The overall objectives of the restructuring activities were to lower costs and better utilize our overall existing manufacturing capacity. During fiscal 2010, we continued this effort to further increase operating efficiencies although we implemented fewer changes than in the prior three fiscal years. Our efforts have helped enhance our ability to improve operating margins, retain and expand existing relationships with customers and attract new business. We may utilize similar measures in the future to realign our operations to further increase our operating efficiencies. The effect of these efforts may materially affect our future operating results.

Fiscal 2010 Compared with Fiscal 2009. During fiscal 2010, we incurred \$2.9 million of restructuring and other charges related to headcount reductions and facility closures. Of this \$2.9 million of restructuring costs, \$1.3 million was recorded within our Healthcare division, \$0.5 million within our Security division, \$1.0 million within our Optoelectronics and Manufacturing division and \$0.1 million in our Corporate segment. During fiscal 2009, we incurred total restructuring and other charges of \$7.1 million related to headcount reductions, costs associated with the closure of certain facilities and a non-recurring litigation charge.

Fiscal 2009 Compared with Fiscal 2008. During fiscal 2009, we incurred \$7.1 million of restructuring and other charges. Of this amount, \$2 million related to a non-recurring litigation charge. The remaining \$5.1 million was primarily related to headcount reductions and facility closures. Of this \$5.1 million of restructuring costs, we recorded \$3.3 million within our Healthcare division, \$1.3 million within our Security division, \$0.2 million within our Optoelectronics and Manufacturing division and \$0.3 million in our Corporate segment. The total restructuring and other charges of \$7.1 million compared to restructuring and charges of \$4.7 million in fiscal 2008, related primarily to headcount reductions and costs associated with the closure of certain facilities.

Other Income and Expenses

	<u>2008</u>	<u>% of Net Sales</u>	<u>2009</u> <u>(Dollars in millions)</u>	<u>% of Net Sales</u>	<u>2010</u>	<u>% of Net Sales</u>
Interest expense and other income, net	<u>\$4.4</u>	<u>(0.7)%</u>	<u>\$2.9</u>	<u>(0.5)%</u>	<u>\$1.8</u>	<u>(0.3)%</u>

Interest Expense and Other Income, net

Fiscal 2010 Compared to Fiscal 2009. In fiscal 2010, the \$1.1 million reduction in interest expense and other income, net, resulted from lower market-driven interest rates on our outstanding borrowings as well as lower levels of borrowings as a result of strong, positive cash flow, and a reduction in a contingent liability related to an acquisition recorded as other income in fiscal 2010.

Fiscal 2009 Compared to Fiscal 2008. In fiscal 2009, we incurred interest expense and other income, net, of \$2.9 million, compared to \$4.4 million in fiscal 2008. This improvement was attributable to a lower cost of borrowing, as a result of lower market-driven interest rates, and by lower average levels of debt. We incurred lower levels of debt due to \$44.5 million of cash generated by operations in fiscal 2009.

Provision for Income Taxes

The effective tax rate for a particular period varies depending on a number of factors including (i) the mix of income earned in various tax jurisdictions, each of which applies a unique range of income tax rates and income tax credits, (ii) changes in previously established valuation allowances for deferred tax assets (changes are based upon our current analysis of the likelihood that these deferred tax assets will be realized), (iii) the level of non-deductible expenses and (iv) tax holidays granted to certain of our international subsidiaries.

Fiscal 2010 Compared to Fiscal 2009. In fiscal 2010, our income tax expense was \$11.4 million, compared to an income tax expense of \$5.4 million for fiscal 2009. The effective income tax rate for fiscal 2010 was 32.7%, compared to 32.5% for fiscal 2009. Our provision for income taxes is dependent on the mix of income from U.S. and foreign locations due to tax rate differences among such countries as well the impact of permanent taxable differences.

Fiscal 2009 Compared to Fiscal 2008. In fiscal 2009, our income tax expense was \$5.4 million, compared to an income tax expense of \$0.6 million for fiscal 2008. The effective income tax rate for fiscal 2009 was 32.5%, compared to 4.0% for fiscal 2008. Included within the fiscal 2008 tax expense was a net tax benefit of \$3.9 million as a result of discrete items impacting the tax provision, the largest of which was a \$4.3 million tax benefit associated with the repurchase of the minority interest of Spacelabs Healthcare. However, excluding the impact of the discrete tax benefits noted above, the effective tax rate for fiscal 2008 was 31.3%. Our provision for income taxes is dependent on the mix of income from U.S. and foreign locations due to tax rate differences among such countries as well the impact of permanent taxable differences.

Liquidity and Capital Resources

Over the past several years we have financed our business primarily from operations and by utilizing our credit facilities, when necessary. In fiscal 2010, our levels of debt decreased as a result of strong cash flow. The changes in our working capital and cash and cash equivalent balances are described below.

	<u>2008</u>	<u>2009</u>	<u>2010</u>	<u>2008-2009</u>	<u>2009-2010</u>
			(Dollars in millions)	% Change	% Change
Working capital	\$ 195.0	\$ 187.6	\$ 204.6	(4)%	9%
Cash and cash equivalents	18.2	25.2	52.0	38%	106%

Working Capital

Fiscal 2010 Compared to Fiscal 2009. Working capital increased by \$17.0 million, or 9%, during fiscal 2010. The most significant increases in working capital were due to cash and cash equivalents increasing by \$26.8 million, or 106%, to \$52.0 in fiscal 2010 from \$25.2 million in fiscal 2009 and accounts receivable, which increased by \$22.3 million, or 20%, to \$132.7 million in fiscal 2010 from \$110.5 million in fiscal 2009. The increase in cash and cash equivalents mainly reflects the significant generation of operating cash flow

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accomplished in fiscal 2010. The increase in accounts receivable reflects the 19% increase in revenue in the three months ended June 30, 2010 as compared to the comparable prior year period. In addition, working capital increased in fiscal 2010 due to a \$5.3 million decrease in accounts payable. The most significant decreases to working capital were (i) a \$24.8 million decrease in inventory, as a result of inventory reduction initiatives in all three divisions and (ii) a \$12.5 million increase in advances by customers, primarily in our Security division.

Fiscal 2009 Compared to Fiscal 2008. Working capital decreased by \$7.4 million, or 4%, during fiscal 2009. The most significant decrease in working capital was accounts receivable, which decreased by \$46.3 million, or 30%, to \$110.5 million in fiscal 2009 from \$156.9 million in fiscal 2008. This decrease was a result of an effective, ongoing focus on collections as well as a 5% reduction in revenue in fiscal 2009 as compared to fiscal 2008. The most significant increases to working capital were (i) \$11.1 million of net earnings, (ii) a \$20.3 million decrease in account payable and (iii) net repayments of bank lines of credit of \$14.6 million during fiscal 2009. Such repayments were a result of the positive cash flow generated by our improved earnings and improved management of working capital.

	<u>2008</u>	<u>2009</u>	<u>2010</u>	<u>2008-2009</u>	<u>2009-2010</u>
			(Dollars in millions)	% Change	% Change
Cash provided by (used in):					
Operating activities	\$ (0.7)	\$ 44.5	\$ 52.1	NM	17%
Investing activities	(29.8)	(12.4)	(24.4)	(58)%	97%
Financing activities	33.0	(24.8)	(3.0)	(175)%	(88)%

Cash Provided by (Used) in Operating Activities

Cash flows from operating activities can fluctuate significantly from period to period as profitability, tax timing differences and other items can significantly impact cash flows. Our largest source of operating cash flows is cash collections from our customers following the sale of our products and services. Our primary uses of cash for operating activities are for purchasing inventory in support of the products that we sell, personnel related expenditures, facilities costs and payments for general operating matters.

Fiscal 2010 Compared to Fiscal 2009. Cash generated by operating activities in fiscal 2010 was \$52.1 million, an increase of \$7.6 million, or 17%, from fiscal 2009, when our operating activities generated \$44.5 million of cash. This improvement was partially due to a \$14.7 million increase in our net income in fiscal 2010 as compared to fiscal 2009, after giving consideration to non-cash operating items, including depreciation and amortization, stock-based compensation, deferred taxes and provisions for losses on accounts receivable, among others for both periods. This improvement was also due to increased emphasis on better working capital management during fiscal 2010, resulting in: (i) a \$37.5 million reduction in the change in inventory, due primarily to our inventory reduction initiatives, (ii) a \$9.9 million increase in the change from accounts payables, (iii) a \$5.0 million reduction in the change in other receivables and (iv) a \$4.3 million increase in the change in advances from customers.

These cash generating improvements were partially offset by: (i) an increase in the change in account receivables of \$54.2 million, partially due to the 19% increase in revenue in the three months ended June 30 2010 as compared to the comparable prior year period and (ii) a decrease in the change in other accrued expenses and current liabilities of \$7.7 million.

Fiscal 2009 Compared to Fiscal 2008. Cash generated by operating activities in fiscal 2009 was \$44.5 million, an increase of \$45.2 million from fiscal 2008, when our operating activities used \$0.7 million of cash. This improvement was partially due to a \$3.3 million increase in our net income in fiscal 2009 as compared to fiscal 2008, after giving consideration to non-cash operating items, including depreciation and amortization, stock-based compensation, deferred taxes and provisions for losses on accounts receivable, among others for both periods. This improvement was also due to increased emphasis on better working capital management

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during fiscal 2009, resulting in: (i) a \$45.9 million reduction in the change in accounts receivables, due primarily to an ongoing focus on collections, (ii) a \$19.9 million increase in change from advances from customers, primarily reflecting significant large sales contracts entered into by our Security division during fiscal 2009, (iii) an \$8.0 million decrease in the change in inventory and (iv) an increase in the change in deferred revenue of \$4.0 million. These cash generating improvements were partially offset by: (i) a decrease in the change in accounts payable of \$29.1 million, (ii) an increase in the change in other receivable of \$4.7 million and (iii) decrease in the change in accrued warranty of \$4.7 million.

Cash Used in Investing Activities

The changes in cash flows from investing activities primarily related to capital expenditures as well as an acquisition of a business and other assets to support our growth plans.

Fiscal 2010 Compared to Fiscal 2009. Net cash used in investing activities was \$24.4 million in fiscal 2010, an increase of \$12.0 million, or 97%, as compared to the \$12.4 million used in fiscal 2009. Such increase was primarily due to the manufacture of cargo screening systems destined for use in connection with the provision of turn-key cargo screening services for the Puerto Rico Port Authority. During fiscal 2010, we also paid \$3.2 million related to an acquisition. There were no acquisitions in the prior-year period.

Fiscal 2009 Compared to Fiscal 2008. Net cash used in investing activities was \$12.4 million in fiscal 2009, a decrease of \$17.4 million in cash used as compared to the \$29.8 million used in investing activities in fiscal 2008. In fiscal 2009, the primary investing activity involved \$10.9 million of capital expenditures as compared to \$12.1 million during the comparable prior year period. In addition, in fiscal 2008 we used \$15.7 million of cash to repurchase shares of Spacelabs Healthcare stock with no comparable investing activity in fiscal 2009. Partially offsetting this reduced investing use of cash in fiscal 2009 as compared to fiscal 2008 was the acquisition of intangible and other assets of \$3.5 million in fiscal 2009, as compared to \$2.5 million in fiscal 2008.

Cash Provided by Financing Activities

The changes in cash flows from financing activities primarily relate to: (i) borrowings and payments under debt obligations, (ii) the issuance of and/or repurchase of common stock and (iii) the exercise activity in our equity participation and employee stock purchase plans.

Fiscal 2010 Compared to Fiscal 2009. Net cash used in financing activities was \$3.0 million in fiscal 2010, a \$21.8 million, or 88%, decrease as compared to net cash of \$24.8 million used in fiscal 2009. During fiscal 2010, we used \$12.0 million in cash to pay down our ongoing scheduled debt and capital leases and \$4.0 million to pay off our bank lines of credit. In fiscal 2009, we paid down \$6.6 million of scheduled debt and capital leases and \$14.4 million to reduce our bank lines of credit. In fiscal 2010 we received proceeds of \$13.0 million from the exercise of stock options and employee stock purchase plan as compared to \$3.6 million in fiscal 2009. In addition, in fiscal 2010 we did not repurchase any shares of our Common Stock under the ongoing authorized Common Stock repurchase program. During fiscal 2009, we used \$7.4 million in cash to repurchase 619,768 shares of our Common Stock.

Fiscal 2009 Compared to Fiscal 2008. Net cash used in financing activities was \$24.8 million in fiscal 2009, a \$57.8 million decrease as compared to net cash provided by financing activity of \$33.0 million in fiscal 2008. During fiscal 2009, we used \$7.4 million in cash to repurchase 619,768 shares of our Common Stock. We also paid down our ongoing scheduled debt and capital leases by \$6.6 million and our revolving lines of credit by \$14.4 million. In fiscal 2008, we received net proceeds of \$23.8 million when we entered into a new credit agreement while simultaneously paying down the preceding credit facility, less the ongoing repayment of our new credit agreement as well as all other scheduled debt and capital lease payments. In addition, in fiscal 2008 we received net proceeds of \$1.9 million from our revolving lines of credit.

Borrowings

We maintain a credit agreement with certain lenders allowing for initial borrowings of up to \$124.5 million. The credit agreement consists of a \$74.5 million, five-year, revolving credit facility (including a \$64.5 million sub-limit for letters-of-credit) and a \$50 million five-year term loan. Borrowings under the agreement bear interest at either (i) the London Interbank Offered Rate (LIBOR) plus between 2.00% and 2.50% or (ii) the bank’s prime rate plus between 1.00% and 1.50%. The rates are determined based on our consolidated leverage ratio. As of June 30, 2010, the weighted-average interest rate under the credit agreement was 2.35%. Our borrowings under the credit agreement are guaranteed by our domestic subsidiaries and are secured by substantially all of our subsidiary guarantors’ assets. The agreement contains various representations, warranties, affirmative, negative and financial covenants, and conditions of default customary for financing agreements of this type. As of June 30, 2010, \$32.3 million was outstanding under the term loan, while no debt was outstanding under the revolving credit facility and \$38.4 million was outstanding under the letter-of-credit facility.

Several of our foreign subsidiaries maintain bank lines-of-credit, denominated in local currencies, to meet short-term working capital requirements and for the issuance of letters-of-credit. As of June 30, 2010, \$10.0 million was outstanding under these letter-of-credit facilities, while no debt was outstanding. As of June 30, 2010, the total amount available under these credit facilities was \$18.0 million, with a total cash borrowing sub-limit of \$4.0 million.

In fiscal 2005, we entered into a bank loan of \$5.3 million to fund the acquisition of land and buildings in the U.K. The loan is payable over a 20-year period. The loan bears interest at British pound-based LIBOR plus 1.2%, payable on a quarterly basis. As of June 30, 2010, \$3.0 million remained outstanding under this loan at an interest rate of 1.9% per annum.

The following is a summary of our contractual obligations and commitments at June 30, 2010 (in thousands):

	Payments Due by Period				
	Total	Less than 1 year	2-3 years	4-5 years	After 5 years
Total debt (excluding capital lease obligations)	\$ 35,398	\$ 11,920	\$ 21,015	\$ 436	\$ 2,027
Capital lease obligations	\$ 711	\$ 711	\$ —	\$ —	\$ —
Operating leases	\$ 38,197	\$ 10,794	\$ 17,197	\$ 9,749	\$ 457
Purchase obligations	\$ 61,526	\$ 60,994	\$ 306	\$ 226	\$ —
Defined benefit plan obligation	\$ 7,921	\$ 879	\$ 550	\$ 514	\$ 5,978
Total contractual obligations	<u>\$ 143,753</u>	<u>\$ 85,298</u>	<u>\$ 39,068</u>	<u>\$ 10,925</u>	<u>\$ 8,462</u>
Other Commercial Commitments—letters of credit	<u>\$ 48,370</u>	<u>\$ 41,331</u>	<u>\$ 5,693</u>	<u>\$ 126</u>	<u>\$ 1,220</u>

We anticipate that cash generated from our operations, in addition to existing cash borrowing arrangements and future access to capital markets should be sufficient to meet our cash requirements for the foreseeable future. However, our future capital requirements will depend on many factors, including future business acquisitions, litigation, stock repurchases and levels of research and development spending, among other factors and the adequacy of available funds will depend on many factors, including the success of our businesses in generating cash, continued compliance with financial covenants contained in our credit facility, and the capital markets in general, among other factors.

Stock Repurchase Program

Our Board of Directors has authorized a stock repurchase program under which we may repurchase up to 3,000,000 shares of our Common Stock. During fiscal 2009, we repurchased 619,768 shares under this program, but we repurchased no shares during fiscal 2010. As of June 30, 2010, 711,205 shares were available for

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additional repurchase under the program. Upon repurchase, the shares were restored to the status of authorized but unissued shares and we recorded them as a reduction in the number of shares of Common Stock issued and outstanding in our Consolidated Financial Statements.

Off Balance Sheet Arrangements

As of June 30, 2010, we had no off balance sheet arrangements, as defined in Item 303(a)(4) of Regulation S-K, other than those previously disclosed.

New Accounting Pronouncements

For information with respect to new accounting pronouncements and the impact of these pronouncements on our Consolidated Financial Statements, see Note 1 to Consolidated Financial Statements.

Related-Party Transactions

In 1994, we, together with an unrelated company, formed ECIL-Rapiscan Security Products Limited, a joint venture organized under the laws of India. We own a 36% interest in the joint venture, our Chairman and Chief Executive Officer owns a 10.5% interest, and our Executive Vice President and the President of our Security division owns a 4.5% ownership interest. Our initial investment was \$0.1 million. For the years ended June 30, 2008, 2009 and 2010, our equity earnings in the joint venture amounted to \$0.4 million, \$0.5 million and \$0.4 million, respectively. We, our Chairman and Chief Executive Officer and our Executive Vice President and the President of our Security division collectively control less than 50% of the board of directors voting power in the joint venture. As a result, we account for the investment under the equity method of accounting. The joint venture was formed for the purpose of the manufacture, assembly, service and testing of security and inspection systems and other products. Some of our subsidiaries are suppliers to the joint venture, which in turn manufactures and sells the resulting products. Sales to the joint venture for fiscal 2008, 2009, and 2010 were approximately \$1.6 million, \$4.4 million and \$4.4 million, respectively. Receivables from the joint venture were \$2.7 million and \$0.9 million as of June 30, 2009 and 2010, respectively.

We have contracted with entities owned by members of our Board of Directors and/or their family members to provide messenger services, auto rental and printing services. Included in cost of sales and selling, general and administrative expenses for the fiscal 2008, 2009, and 2010, are approximately \$40,000, \$54,000 and \$64,000, respectively, for messenger service and auto rental; and \$42,000, \$45,000 and \$60,000, respectively, for printing services.

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UNAUDITED QUARTERLY RESULTS

The following tables present unaudited quarterly financial information for the four quarters ended June 30, 2009 and 2010 (in thousands, except per share data):

	Quarter Ended			
	September 30, 2008	December 31, 2008	March 31, 2009	June 30, 2009
	(Unaudited)			
Revenues	\$ 148,161	\$ 159,042	\$ 144,095	\$ 139,063
Costs of goods sold	98,526	104,623	94,264	91,497
Gross profit	49,635	54,419	49,831	47,566
Operating expenses:				
Selling, general and administrative expenses	37,541	35,727	34,406	30,311
Research and development	10,213	8,669	8,572	9,408
Restructuring and other charges	801	2,798	2,401	1,123
Total operating expenses	48,555	47,194	45,379	40,842
Income from operations	1,080	7,225	4,452	6,724
Interest expense and other income, net	895	863	583	595
Income before provision for income taxes	185	6,362	3,869	6,129
Provision for income taxes	53	2,200	1,296	1,844
Net income	\$ 132	\$ 4,162	\$ 2,573	\$ 4,285
Basic earnings per common share	\$ 0.01	\$ 0.24	\$ 0.15	\$ 0.25
Diluted earnings per common share	\$ 0.01	\$ 0.24	\$ 0.15	\$ 0.24

	Quarter Ended			
	September 30, 2009	December 31, 2009	March 31, 2010	June 30, 2010
	(Unaudited)			
Revenues	\$ 133,761	\$ 150,621	\$ 145,401	\$ 165,328
Costs of goods sold	89,294	94,256	92,184	101,343
Gross profit	44,467	56,365	53,217	63,985
Operating expenses:				
Selling, general and administrative expenses	32,227	34,610	34,789	38,204
Research and development	7,989	10,353	9,129	11,106
Restructuring and other charges	53	607	946	1,253
Total operating expenses	40,269	45,570	44,864	50,563
Income from operations	4,198	10,795	8,353	13,422
Interest expense and other income, net	605	784	(175)	558
Income before provision for income taxes	3,593	10,011	8,528	12,864
Provision for income taxes	1,083	3,059	2,416	4,881
Net income	\$ 2,510	\$ 6,952	\$ 6,112	\$ 7,983
Basic earnings per common share	\$ 0.14	\$ 0.39	\$ 0.34	\$ 0.44
Diluted earnings per common share	\$ 0.14	\$ 0.39	\$ 0.33	\$ 0.42

ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

Market Risk

We are exposed to certain market risks, which are inherent in our financial instruments and arise from transactions entered into in the normal course of business. We may enter into derivative financial instrument transactions in order to manage or reduce market risk in connection with specific foreign-currency-denominated transactions. We do not enter into derivative financial instrument transactions for speculative purposes.

We are subject to interest rate risk on our short-term borrowings under our bank lines of credit. Borrowings under these lines of credit do not give rise to significant interest rate risk because these borrowings have short maturities and are borrowed at variable interest rates. Historically, we have not experienced material gains or losses due to interest rate changes.

Foreign Currency

We maintain the accounts of our operations in each of the following countries in the respective currencies: Finland, France, Germany, Italy and Greece (Euros), Singapore (U.S. dollars), Malaysia (Malaysian ringgits), United Kingdom (U.K. pounds), Norway (Norwegian kroners), India (Indian rupees), Indonesia (Indonesian rupiah and U.S. dollars), Hong Kong (Hong Kong dollars), China (Chinese Yuan), Canada (Canadian dollars), Mexico (Mexican pesos), Australia (Australian dollars) and Cyprus (Cypriot pounds). Foreign currency financial statements are translated into U.S. dollars at fiscal year-end rates, with the exception of revenues, costs and expenses, which are translated at average rates during the reporting period. We include gains and losses resulting from foreign currency transactions in income, while we exclude those resulting from translation of financial statements from income and include them as a component of accumulated other comprehensive income. Transaction gains and losses, which were included in our consolidated statement of operations, amounted to a gain of approximately \$0.8 million, a gain of \$0.4 million and a loss of \$3.2 million for the fiscal years ended June 30, 2008, 2009 and 2010, respectively. Furthermore, a 10% appreciation of the U.S. dollar relative to the local currency exchange rates would have resulted in a net increase in our operating income of approximately \$7.0 million in fiscal 2010. Conversely, a 10% depreciation of the U.S. dollar relative to the local currency exchange rates would have resulted in a net decrease in our operating income of approximately \$7.0 million in fiscal 2010.

Use of Derivatives

Our use of derivatives consists primarily of foreign exchange contracts and interest rate swap agreements. As discussed in Note 1 to the Consolidated Financial Statements, as of June 30, 2010, we had outstanding an interest rate swap agreement, which was considered effective cash flow hedges in their entirety. As a result, the net loss on such derivative contract has been reported as a component of other comprehensive income in the Consolidated Financial Statements and will be reclassified into net earnings when the hedged transaction settles.

Importance of International Markets

International markets provide us with significant growth opportunities. However, the following events, among others, could adversely affect our financial results in subsequent periods: periodic economic downturns in different regions of the world, changes in trade policies or tariffs, civil or military conflict and other political instability. We continue to perform ongoing credit evaluations of our customers' financial condition and, if deemed necessary, we require advance payments for sales. We monitor economic and currency conditions around the world to evaluate whether there may be any significant effect on our international sales in the future. Due to our overseas investments and the necessity of dealing with local currencies in our foreign business transactions, we are at risk with respect to foreign currency fluctuations.

Inflation

We do not believe that inflation has had a material impact on our results of operations.

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Interest Rate Risk

We utilize short-term and long-term financing and may use interest rate hedges to manage the effect of interest rate changes on our existing debt. As of June 30, 2010, we had an interest rate swap agreement outstanding as discussed above under "Use of Derivatives."

The principal maturity and estimated value of our long-term debt exposure as of June 30, 2009 are as follows (in thousands):

	Maturity						Total	Fair Value
	2010	2011	2012	2013	2014	2015 and thereafter		
Secured long term loans and capital lease obligations	\$ 8,577	\$ 11,007	\$ 7,798	\$ 18,042	\$ 273	\$ 2,663	\$ 48,360	\$ 48,360
Average interest rate	3.2%	3.5%	4.3%	3.9%	4.0%	4.0%	3.2%	

The principal maturity and estimated value of our long-term debt exposure as of June 30, 2010 are as follows (in thousands):

	Maturity						Total	Fair Value
	2011	2012	2013	2014	2011	2016 and thereafter		
Secured long term loans and capital lease obligations	\$ 12,743	\$ 6,294	\$ 14,609	\$ 218	\$ 218	\$ 2,027	\$ 36,109	\$ 36,109
Average interest rate	3.3%	3.5%	2.9%	3.4%	3.4%	3.4%	3.3%	

ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA

We make reference here to the Index to Consolidated Financial Statements that appears on page F-1 of this report. The Report of Independent Registered Public Accounting Firm from Moss Adams LLP, the Consolidated Financial Statements and the Notes to Consolidated Financial Statements listed in the Index to Consolidated Financial Statements, which appear beginning on page F-2 of this report, are incorporated by reference into this Item 8.

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

As of June 30, 2010, the end of the period covered by this report, our management, including our Chief Executive Officer and our Chief Financial Officer, reviewed and evaluated the effectiveness of our disclosure controls and procedures (as defined in Rule 13a-15(e) and 15d-15(e) of the Securities Exchange Act of 1934, as amended). Such disclosure controls and procedures are designed to ensure that material information we must disclose in this report is recorded, processed, summarized and filed or submitted on a timely basis. Based upon that evaluation our management, Chief Executive Officer and Chief Financial Officer, concluded that our disclosure controls and procedures were effective as of June 30, 2010.

Management’s Report on Internal Control over Financial Reporting

Our management is responsible for establishing and maintaining adequate internal control over financial reporting, as such term is defined in Exchange Act Rule 13a-15(f). Under the supervision and with the participation of management, including the Chief Executive Officer and our Chief Financial Officer, we conducted an evaluation of the effectiveness of our internal control over financial reporting based on the framework in *Internal Control—Integrated Framework* issued by the Committee of Sponsoring Organizations of the Treadway Commission (COSO). Based on that evaluation, management concluded that our internal control over financial reporting was effective as of June 30, 2010.

Moss Adams LLP, an independent registered public accounting firm that audited the financial statements included in this report, has issued its attestation report, which appears below, on the effectiveness of our internal controls over financial reporting.

Changes in Internal Control over Financial Reporting

There were no changes in our internal control over financial reporting during fiscal 2010 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

Report of Independent Registered Public Accounting Firm

To the Board of Directors and Shareholders of OSI Systems, Inc.:
Hawthorne, California

We have audited OSI Systems, Inc. and subsidiaries, (the Company) internal control over financial reporting as of June 30, 2010, based on criteria established by the Committee of Sponsoring Organizations of the Treadway Commission (COSO) in Internal Control—Integrated Framework. The Company’s management is responsible for maintaining effective internal control over financial reporting and for its assessment of the effectiveness of internal control over financial reporting. 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 includes performing such other procedures, as we consider 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 accounting principles generally accepted in the United States of America. 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 consolidated 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, OSI Systems, Inc. and subsidiaries maintained, in all material respects, effective internal control over financial reporting as of June 30, 2010, based on the COSO criteria.

We have also audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States), the consolidated financial statements and financial statement schedule of OSI Systems, Inc. and subsidiaries as of and for the year ended June 30, 2010, and our report dated August 26, 2010 expressed an unqualified opinion on those consolidated financial statements and financial statement schedule.

/s/ MOSS ADAMS LLP

Los Angeles, California
August 26, 2010

ITEM 9B. OTHER INFORMATION

During fiscal 2010, we reincorporated our company from California to Delaware. In connection with this reincorporation, we modified the form of indemnification agreement that members of our Board of Directors and certain of our executive officers sign. We did so in order to reflect in the indemnification agreement the fact that our company is now incorporated in Delaware. We have filed a copy of the new form of indemnification agreement as Exhibit 10.17 to this Annual Report. As of the date of this Annual Report, all of our directors and executive officer have either entered into this new form indemnification agreement or have amended their current indemnification agreement such that they are now bound, in effect, to the terms of the new form.

PART III

ITEM 10. DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE

The information called for by this item is hereby incorporated by reference from our definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders, which Proxy Statement is anticipated to be filed with the Securities and Exchange Commission within 120 days of June 30, 2010.

ITEM 11. EXECUTIVE COMPENSATION

The information called for by this item is hereby incorporated by reference from our definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders, which Proxy Statement is anticipated to be filed with the Securities and Exchange Commission within 120 days of June 30, 2010.

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

The information called for by this item is hereby incorporated by reference from our definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders, which Proxy Statement is anticipated to be filed with the Securities and Exchange Commission within 120 days of June 30, 2010.

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

The information called for by this item is hereby incorporated by reference from our definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders, which Proxy Statement is anticipated to be filed with the Securities and Exchange Commission within 120 days of June 30, 2010.

ITEM 14. PRINCIPAL ACCOUNTING FEES AND SERVICES

The information called for by this item is hereby incorporated by reference from our definitive Proxy Statement relating to the 2010 Annual Meeting of Shareholders, which Proxy Statement is anticipated to be filed with the Securities and Exchange Commission within 120 days of June 30, 2010.

PART IV

ITEM 15. EXHIBITS, FINANCIAL STATEMENT SCHEDULES

(a) The following documents are filed as part of this report:

1. *Financial Statements.* Please see the accompanying Index to Consolidated Financial Statements, which appears on page F-1 of the report. The Report of Independent Registered Public Accounting Firm, the Consolidated Financial Statements and the Notes to Consolidated Financial Statements listed in the Index to Consolidated Financial Statements, which appear beginning on page F-2 of this report, are incorporated by reference into Item 8 above.

2. *Financial Statement Schedules.*

Schedule II—Valuation and Qualifying Accounts

No other financial statement schedules are presented as the required information is either not applicable or included in the Consolidated Financial Statements or notes thereto.

(b) *Exhibits.* The exhibits listed on the accompanying Exhibit Index immediately following the signature page are filed as part of, or are incorporated by reference into, this report.

(c) *Financial Statement Schedules.* Reference is made to Item 15(a)(2) above.

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OSI SYSTEMS, INC.
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REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Board of Directors and Shareholders of OSI Systems, Inc.:

We have audited the accompanying consolidated balance sheets of OSI Systems, Inc. and Subsidiaries as of June 30, 2009 and 2010, and the related consolidated statements of operations, shareholders' equity and cash flows for the three years ended June 30, 2008, 2009 and 2010. Our audits also included the financial statement schedule listed in the index at Item 15 in Schedule II. These financial statements and financial statement schedule are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements and financial statement schedule 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 consolidated financial statements are free of material misstatement. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the consolidated financial statements, 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 consolidated financial position of OSI Systems, Inc. and Subsidiaries as of June 30, 2009 and 2010, and the consolidated results of its operations and cash flows for the years ended June 30, 2008, 2009 and 2010, in conformity with accounting principles generally accepted in the United States of America. Also, in our opinion, such financial statement schedule, when considered in relation to the basic consolidated financial statements taken as a whole, presents fairly, in all material respects, the information set forth therein.

As discussed in Note 14 to the consolidated financial statements, the Company adopted Statement of Financial Accounting Standard No. 158 "Employers' Accounting for Defined Benefit Pension and Other Postretirement Plans—an Amendment of FASB Statements No. 87, 88, 106 and 132(R)," (codified in FASB ASC Topic 715, Compensation—Retirement Plans) which changed the Company's method of accounting for pension and postretirement benefits as of June 30, 2007. As discussed in Notes 1 and 10 to the consolidated financial statements, effective July 1, 2007, the Company adopted FASB Interpretation No. 48, "Accounting for Uncertainty in Income Taxes—an Interpretation of FASB No. 109" (codified in FASB ASC Topic 740, Income Taxes).

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States), the effectiveness of OSI Systems, Inc. and Subsidiaries' internal control over financial reporting as of June 30, 2010, based on criteria established in Internal Control—Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission, and our report dated August 26, 2010 expressed an unqualified opinion thereon.

/s/ MOSS ADAMS LLP
Los Angeles, California
August 26, 2010

OSI SYSTEMS, INC. AND SUBSIDIARIES
CONSOLIDATED BALANCE SHEETS
(in thousands, except share data)

	<u>June 30,</u>	
	<u>2009</u>	<u>2010</u>
ASSETS		
CURRENT ASSETS:		
Cash and cash equivalents	\$ 25,172	\$ 51,989
Accounts receivable	110,453	132,728
Other receivables	2,950	2,859
Inventories	150,763	125,930
Deferred income taxes	20,128	17,262
Prepaid expenses and other current assets	13,777	18,433
Total current assets	<u>323,243</u>	<u>349,201</u>
Property and equipment, net	42,232	51,515
Goodwill	60,195	63,941
Intangible assets, net	32,451	31,975
Other assets	16,707	16,482
Total assets	<u>\$474,828</u>	<u>\$513,114</u>
LIABILITIES AND SHAREHOLDERS' EQUITY		
CURRENT LIABILITIES:		
Bank lines of credit	\$ 4,000	\$ —
Current portion of long-term debt	8,557	12,743
Accounts payable	54,980	49,673
Accrued payroll and related expenses	22,416	23,953
Advances from customers	12,863	25,325
Accrued warranties	10,106	10,930
Deferred revenue	8,880	7,698
Other accrued expenses and current liabilities	13,833	14,272
Total current liabilities	<u>135,635</u>	<u>144,594</u>
Long-term debt	39,803	23,366
Other long-term liabilities	23,390	31,444
Total liabilities	<u>198,828</u>	<u>199,404</u>
Commitment and contingencies (Note 11)		
Shareholders' Equity:		
Preferred stock, no par value—authorized, 10,000,000 shares; no shares issued or outstanding	—	—
Common stock, \$0.001 par value—authorized, 100,000,000 shares; issued and outstanding, 17,411,569 and 18,326,133 shares at June 30, 2009 and 2010, respectively	225,297	244,026
Retained earnings	53,124	76,681
Accumulated other comprehensive loss	(2,421)	(6,997)
Total shareholders' equity	<u>276,000</u>	<u>313,710</u>
Total liabilities and shareholders' equity	<u>\$474,828</u>	<u>\$513,114</u>

See accompanying notes to Consolidated Financial Statements.

OSI SYSTEMS, INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF OPERATIONS
(in thousands, except per share data)

	Year Ended June 30,		
	2008	2009	2010
Revenues	\$ 623,088	\$ 590,361	\$ 595,111
Cost of goods sold	404,049	388,910	377,077
Gross profit	<u>219,039</u>	<u>201,451</u>	<u>218,034</u>
Operating expenses:			
Selling, general and administrative expenses	150,082	137,985	139,830
Research and development	45,361	36,862	38,577
Restructuring and other charges	4,688	7,123	2,859
Total operating expenses	<u>200,131</u>	<u>181,970</u>	<u>181,266</u>
Income from operations	18,908	19,481	36,768
Interest expense and other income, net	4,469	2,936	1,772
Income before income taxes	14,439	16,545	34,996
Provision for income taxes	579	5,393	11,439
Net income	<u>\$ 13,860</u>	<u>\$ 11,152</u>	<u>\$ 23,557</u>
Earnings per share:			
Basic	<u>\$ 0.80</u>	<u>\$ 0.64</u>	<u>\$ 1.32</u>
Diluted	<u>\$ 0.78</u>	<u>\$ 0.63</u>	<u>\$ 1.28</u>
Shares used in per share calculation:			
Basic	<u>17,428</u>	<u>17,518</u>	<u>17,874</u>
Diluted	<u>17,735</u>	<u>17,596</u>	<u>18,389</u>

See accompanying notes to Consolidated Financial Statements.

OSI SYSTEMS, INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF SHAREHOLDERS' EQUITY
(in thousands, except share data)

	Common		Retained Earnings	Accumulated Other Comprehensive (loss) Income	Comprehensive (loss) Income	Total
	Number of Shares	Amount				
Balance—June 30, 2007	17,086,989	\$207,260	\$ 31,450	\$ 8,502		\$247,212
Exercise of stock options	340,642	5,807				5,807
Vesting of restricted shares	6,671					—
Net tax expense of stock options exercised/cancelled		(411)				(411)
Shares purchased under employee stock purchase program	65,908	1,296				1,296
Stock compensation expense		4,777				4,777
Shares issued for the purchase of Spacelabs Healthcare shares	239,847	5,898				5,898
Dividend distribution to Minority Interest Shareholders		(46)				(46)
FIN 48 tax adjustment			(3,338)			(3,338)
Comprehensive income (loss):						
Net income			13,860			13,860
Translation adjustment				3,276		3,276
Minimum pension liability adjustment—net of tax				(310)		(310)
Comprehensive income					\$ 16,826	
Balance—June 30, 2008	17,740,057	\$224,581	\$ 41,972	\$ 11,468		\$278,021
Exercise of stock options	163,680	2,495				2,495
Vesting of restricted shares	52,006					—
Net tax expense of stock options exercised/cancelled		(555)				(555)
Shares purchased under employee stock purchase program	75,594	1,109				1,109
Stock compensation expense		5,055				5,055
Treasury shares	(619,768)	(7,388)				(7,388)
Comprehensive income (loss):						
Net income			11,152			11,152
Translation adjustment				(13,644)		(13,644)
Minimum pension liability adjustment—net of tax				(529)		(529)
Other				284		284
Comprehensive loss					\$ (2,737)	
Balance—June 30, 2009	17,411,569	\$225,297	\$ 53,124	\$ (2,421)		\$276,000
Exercise of stock options	660,764	11,226				11,226
Vesting of restricted shares	112,664					—
Net tax benefit of stock options exercised/cancelled		732				732
Shares purchased under employee stock purchase program	141,136	1,760				1,760
Stock compensation expense		5,011				5,011
Comprehensive income (loss):						
Net income			23,557			23,557
Translation adjustment				(3,202)		(3,202)
Minimum pension liability adjustment—net of tax				(296)		(296)
Net unrealized loss on investments				(2,097)		(2,097)
Net tax benefit from unrealized loss on investments and derivative instrument				693		693
Other				326		326
Comprehensive income					\$ 18,981	
Balance—June 30, 2010	<u>18,326,133</u>	<u>\$244,026</u>	<u>\$ 76,681</u>	<u>\$ (6,997)</u>		<u>\$313,710</u>

See accompanying notes to Consolidated Financial Statements.

OSI SYSTEMS, INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF CASH FLOWS
(in thousands)

	Year Ended June 30,		
	2008	2009	2010
CASH FLOWS FROM OPERATING ACTIVITIES			
Net income	\$ 13,860	\$ 11,152	\$ 23,557
Adjustments to reconcile net income to net cash provided by (used in) operating activities:			
Depreciation and amortization	19,342	17,805	18,561
Stock based compensation expense	4,777	5,055	5,011
Provision for losses on accounts receivable	335	5,377	291
Equity in earnings of unconsolidated affiliates	(403)	(689)	(402)
Tax effect of exercise (cancellation) of stock options	(411)	(555)	732
Deferred income taxes	(4,887)	(2,259)	2,714
Other	156	200	289
Changes in operating assets and liabilities—net of business acquisitions:			
Accounts receivable	(15,127)	30,821	(23,373)
Other receivables	2,009	(2,725)	2,234
Inventories	(25,557)	(17,607)	19,898
Prepaid expenses and other current assets	(3,439)	(1,660)	(4,213)
Accounts payable	13,905	(15,226)	(5,375)
Accrued payroll and related expenses	3,539	667	4,865
Advances from customers	(10,052)	9,806	14,145
Accrued warranties	4,055	(615)	1,065
Deferred revenue	(20)	3,996	(1,087)
Other accrued expenses and current liabilities	(2,754)	957	(6,762)
Net cash provided by (used in) operating activities	<u>(672)</u>	<u>44,500</u>	<u>52,150</u>
CASH FLOWS FROM INVESTING ACTIVITIES			
Acquisition of property and equipment	(12,117)	(10,852)	(18,087)
Proceeds from the sale of property and equipment	488	2,300	37
Acquisition of businesses, net of cash acquired	—	—	(3,241)
Purchase of investments and marketable securities	—	(407)	—
Buyback of subsidiary stock	(15,674)	—	—
Acquisition of intangible and other assets	(2,538)	(3,467)	(3,103)
Net cash used in investing activities	<u>(29,841)</u>	<u>(12,426)</u>	<u>(24,394)</u>
CASH FLOWS FROM FINANCING ACTIVITIES			
Net proceeds from (repayments of) bank lines of credit	1,905	(14,411)	(4,000)
Proceeds from long-term debt	50,127	—	—
Payments on long-term debt	(25,140)	(5,692)	(11,349)
Payments of capital lease obligations	(1,138)	(899)	(644)
Proceeds from exercise of stock options and employee stock purchase plan	7,216	3,605	12,986
Repurchase of common shares	—	(7,388)	—
Net cash provided by (used in) financing activities	<u>32,970</u>	<u>(24,785)</u>	<u>(3,007)</u>
Effect of exchange rate changes on cash	(205)	(349)	2,068
Net increase in cash and cash equivalents	2,252	6,940	26,817
Cash and cash equivalents—beginning of year	15,980	18,232	25,172
Cash and cash equivalents—end of year	<u>\$ 18,232</u>	<u>\$ 25,172</u>	<u>\$ 51,989</u>
Supplemental disclosure of cash flow information:			
Interest	4,411	3,001	2,758
Income taxes	3,229	6,801	7,588
Supplemental disclosure of non-cash investing activities:			
Equipment purchased under capital lease obligations	350	—	—
Buyback of subsidiary stock with Common Stock	5,898	—	—

See accompanying notes to Consolidated Financial Statements.

OSI SYSTEMS, INC. AND SUBSIDIARIES
NOTES TO CONSOLIDATED FINANCIAL STATEMENTS
FOR THE THREE YEARS ENDED JUNE 30, 2010

1. SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

Description of Business—OSI Systems, Inc., together with its subsidiaries (the “Company”), is a vertically integrated designer and manufacturer of specialized electronic systems and components for critical applications. The Company sells its products in diversified markets, including homeland security, healthcare, defense and aerospace.

The Company has three operating divisions: (i) Security, providing security inspection systems and related services; (ii) Healthcare, providing patient monitoring, diagnostic cardiology and anesthesia systems, and related services; and (iii) Optoelectronics and Manufacturing, providing specialized electronic components for the Security and Healthcare divisions as well as for applications in the defense and aerospace markets, among others.

Through its Security division, the Company designs, manufactures, markets and services security and inspection systems worldwide, and provides turnkey security screening services. The Security division’s products are used to inspect baggage, cargo, vehicles and other objects for weapons, explosives, drugs and other contraband and to screen people. These products and services are also used for the safe, accurate and efficient verification of cargo manifests for the purpose of assessing duties and monitoring the export and import of controlled materials.

Through its Healthcare division, the Company designs, manufactures, markets and services patient monitoring, diagnostic cardiology and anesthesia delivery and ventilation systems worldwide primarily under the “Spacelabs” trade name. These products are used by care providers in critical care, emergency and perioperative areas within hospitals as well as physicians’ offices, medical clinics and ambulatory surgery centers.

Through its Optoelectronics and Manufacturing division, the Company designs, manufactures and markets optoelectronic devices and provides electronics manufacturing services worldwide for use in a broad range of applications, including aerospace and defense electronics, security and inspection systems, medical imaging and diagnostics, computed tomography (CT), telecommunications and industrial automation. This division provides products and services to original equipment manufacturers as well as to the Company’s own Security and Healthcare divisions.

Consolidation—The Consolidated Financial Statements include the accounts of OSI Systems, Inc. and its wholly-owned and majority-owned subsidiaries. All significant intercompany accounts and transactions have been eliminated in consolidation. Investments in joint ventures over which the Company has significant influence but does not have voting control are accounted for using the equity method. Investments over which the Company does not have significant influence are accounted for using the cost method.

Spacelabs Healthcare Public Offering and Repurchase—In October 2005, Spacelabs Healthcare, Inc., a subsidiary composed of the business operations of the Company’s Healthcare division, completed an initial public offering of approximately 20% of its total issued and outstanding common stock. The Spacelabs Healthcare shares traded under the ticker symbol “SLAB” on the AIM (formerly known as the Alternative Investment Market), a stock market administered by the London Stock Exchange. During fiscal years 2007 and 2008, the Company repurchased all of the publicly-traded shares of Spacelabs Healthcare, completing this repurchase in the second quarter of fiscal 2008. As a result, the Company owns 100% of the stock of Spacelabs Healthcare.

Use of Estimates—The preparation of financial statements in conformity with accounting principles generally accepted in the United States of America requires management to make estimates and assumptions that

OSI SYSTEMS, INC. AND SUBSIDIARIES
NOTES TO CONSOLIDATED FINANCIAL STATEMENTS—(Continued)
FOR THE THREE YEARS ENDED JUNE 30, 2010

affect the reported amounts of assets and liabilities and the disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period. Actual results could differ from those estimates.

Cash Equivalents—The Company considers all highly liquid investments purchased with maturities of three months or less as of the acquisition date, to be cash equivalents.

Allowance for Doubtful Accounts—The allowance for doubtful accounts involves estimates based on management's judgment, review of individual receivables and analysis of historical bad debts. The Company adjusts customer credit limits based upon each customer's credit worthiness. The Company monitors collections and payments from its customers and maintains allowances for doubtful accounts for estimated losses resulting from the inability of its customers to make required payments. If the financial condition of the Company's customers were to deteriorate, resulting in an impairment of their ability to make payments, additional allowances may be required.

Inventories—Inventories are generally stated at the lower of cost (first-in, first-out) or market. The Company writes down inventory for slow-moving and obsolete inventory based on assessments of future demands, market conditions and customers who may be experiencing financial difficulties. If these factors are less favorable than those projected, additional inventory write-downs may be required.

Unrealized Loss on Investments—The Company owns stock in publically-traded companies, which is classified as available for sale. During fiscal 2010, the market value of these investments declined by \$2.1 million. This unrealized loss is not believed to be other than temporarily impaired and has been charged to other comprehensive income in the consolidated statements of stockholders' equity.

Property and Equipment—Property and equipment are stated at cost less accumulated depreciation and amortization. Depreciation and amortization are computed using the straight-line method over the estimated useful lives of the assets. Amortization of leasehold improvements is calculated on the straight-line basis over the shorter of the useful life of the asset or the lease term. Leased capital assets are included in property and equipment. Amortization of property and equipment under capital leases is included with depreciation expense.

Goodwill and Other Intangible Assets and Valuation of Long-Lived Assets—Goodwill represents the excess purchase price of net tangible and intangible assets acquired in business combinations over their estimated fair value. Goodwill is allocated to the Company's segments based on the nature of the product line of the acquired entity. The carrying value of goodwill is not amortized but is tested for impairment on an annual basis and earlier if there is an indicator of impairment. Intangible assets other than goodwill are amortized over their useful lives unless these lives are determined to be indefinite. The Company tests goodwill for impairment using a two-step process. First, the Company determines if the carrying amount of any of its reporting units, consisting of the Security division, the Healthcare division and the Optoelectronics and Manufacturing division, exceeds its fair value. The fair value of the reporting units was calculated using the income approach and the market approach. Under the income approach, the fair value of the reporting units was calculated by estimating the present value of associated future cash flows. Under the market approach, the fair value was calculated using the guideline public company method and the mergers and acquisitions method. If this testwork indicates a potential impairment of goodwill associated with any reporting unit, the Company then compares the implied fair value of the goodwill associated with the respective reporting unit to its carrying amount to determine if there is an impairment loss. There was no goodwill impairment for fiscal 2008, 2009 and 2010.

The Company evaluates long-lived assets, including intangible assets other than goodwill, for impairment whenever events or changes in circumstances indicate that the carrying amount of the asset may not be

OSI SYSTEMS, INC. AND SUBSIDIARIES
NOTES TO CONSOLIDATED FINANCIAL STATEMENTS—(Continued)
FOR THE THREE YEARS ENDED JUNE 30, 2010

recoverable. An impairment is considered to exist if the total estimated future cash flows on an undiscounted basis are less than the carrying amount of the assets. If impairment does exist, the Company measures the impairment loss and records it based on the discounted estimate of future cash flows. In estimating future cash flows, the Company groups assets that represent the lowest level for which identifiable cash flows are largely independent of the cash flows of other groups of assets and liabilities. The Company's estimate of future cash flows is based upon, among other things, certain assumptions about expected future operating performance, growth rates and other factors.

Income Taxes—Deferred income taxes are provided for temporary differences between the financial statement and income tax basis of the Company's assets and liabilities, based on enacted tax rates. A valuation allowance is provided when it is more likely than not that some portion or all of the deferred income tax assets will not be realized. Income tax accounting standards prescribe a two-step process for the financial statement measurement and recognition of a tax position taken or expected to be taken in a tax return. The first step involves the determination of whether it is more likely than not (greater than 50 percent likelihood) that a tax position will be sustained upon examination, based on the technical merits of the position. The second step requires that any tax position that meets the more-likely-than-not recognition threshold be measured and recognized in the financial statements at the largest amount of benefit that is greater than 50 percent likely of being realized upon ultimate settlement. The income tax accounting standards also provide guidance on the accounting for related interest and penalties, financial statement classification and disclosure. The cumulative effect of applying these standards is to be reported as an adjustment to the opening balance of retained earnings in the period of adoption. See Note 10 for additional information.

Fair Value of Financial Instruments—The Company's financial instruments consist primarily of cash, marketable securities, accounts receivable, accounts payable and debt instruments. The carrying values of financial instruments, other than debt instruments, are representative of their fair values due to their short-term maturities. The carrying values of the Company's long-term debt instruments are considered to approximate their fair values because the interest rates of these instruments are variable or comparable to current rates offered to the Company.

Fair value is the price that would be received to sell an asset or paid to transfer a liability in an orderly transaction between market participants at the measurement date. The Company has determined that all of its marketable securities fall into the "Level 1" category, which values assets at the quoted prices in active markets for identical assets; while the Company's derivative instruments fall into the "Level 2" category, which values assets and liabilities from observable inputs other than quoted market prices. There were no assets or liabilities where "Level 3" valuation techniques were used and there were no assets and liabilities measured at fair value on a non-recurring basis.

The fair values of such assets were:

	<u>June 30,</u>	
	<u>2009</u>	<u>2010</u>
Level 1	\$3,249	\$5,750
Level 2	(342)	(244)
Total	<u>\$2,907</u>	<u>\$5,506</u>

Derivative Instruments and Hedging Activity—The Company's use of derivatives consists primarily of foreign exchange contracts and interest rate swap agreements. To reduce the unpredictability of cash flows from

OSI SYSTEMS, INC. AND SUBSIDIARIES
NOTES TO CONSOLIDATED FINANCIAL STATEMENTS—(Continued)
FOR THE THREE YEARS ENDED JUNE 30, 2010

interest payments related to variable, LIBOR-based debt, as of June 30, 2010 the Company had outstanding a three year interest rate swap agreement, whereby the Company essentially incurs interest expense based upon a fixed 1.69% rate index for a portion of its term loan. The interest rate swap matures in March 2012. Pursuant to Accounting Standards Codification 815 “Derivatives and Hedging,” the interest rate swap was considered an effective cash flow hedge. As a result, the net gains or losses on such derivative contract has been reported as a component of other comprehensive income in the Consolidated Financial Statements and are reclassified into net earnings when the hedged transaction settles.

Revenue Recognition—The Company recognizes revenue upon shipment of products when title and risk of loss passes, and when terms are fixed and collection is probable. The portion of revenue for the sale attributable to installation is deferred and recognized when the installation service is provided. In an instance where terms of sale include subjective customer acceptance criteria, revenue is deferred until the acceptance criteria are met. Concurrent with the shipment of the product, the Company accrues estimated product return reserves and warranty expenses. Critical judgments made by management related to revenue recognition include the determination of whether or not customer acceptance criteria are perfunctory or inconsequential. The determination of whether or not the customer acceptance terms are perfunctory or inconsequential impacts the amount and timing of revenue recognized. Critical judgments also include estimates of warranty reserves, which are established based on historical experience and knowledge of the product.

Revenues from separate service maintenance contracts are recognized ratably over the term of the agreements. For other services, service revenues are recognized as the services are performed. Deferred revenue for services arises from advance payments received from customers for services not yet performed.

Freight—The Company records shipping and handling fees it charges to its customers as revenue and related costs as cost of goods sold.

Research and Development Costs—Research and development costs are those costs related to the development of a new product, process or service, or significant improvement to an existing product, process or service. Such costs are charged to operations as incurred. Grants for research and development are recorded as a reimbursement of costs as such grants are received.

Stock-Based Compensation—Stock-based compensation cost is measured at the grant date based on the estimated fair value of the award and is recognized as expense over the employee’s requisite service period for all stock-based awards granted, modified or cancelled. See Employee Stock Plans at Note 9 to the Consolidated Financial Statements.

Restructuring Charges—The Company consolidates processes and facilities of its subsidiaries to better align with demand by its customers and thereby improve its operational efficiencies. The associated charges, including reducing workforce and capacity, are recognized as restructuring charges in the Consolidated Financial Statements. During fiscal years 2008, 2009 and 2010, the Company consolidated manufacturing processes and facilities of certain businesses that resulted in pre-tax restructuring charges of \$4.7 million, \$7.1 million and \$2.9 million, respectively. See Note 7 for additional information about these restructuring charges.

Concentrations of Credit Risk—Financial instruments that are potentially subject to concentrations of credit risk consist primarily of cash, cash equivalents, marketable securities and accounts receivable. The Company restricts investments in cash equivalents to financial institutions with high credit standing. Credit risk on accounts receivable is minimized as a result of the large and diverse nature of the Company’s worldwide customer base. No individual customer accounted for more than 10% of accounts receivable as of June 30, 2009 or 2010 or 10%

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of revenues for the years ended June 30, 2008, 2009 or 2010. The Company performs ongoing credit evaluations of its customers' financial condition and maintains allowances for potential credit losses. For cost control and efficiency reasons, the Company at times purchases raw materials and subcomponents from a single vendor though it generally qualifies second sources for most of its raw materials and critical components or has identified alternative sources of supply where available.

Foreign Currency Translation—The Company transacts business in various foreign currencies. In general, the functional currency of a foreign operation is the local country's currency. Consequently, revenues and expenses of operations outside the United States are translated into United States dollars using average exchange rates while assets and liabilities of operations outside the United States are translated into United States dollars using year-end exchange rates. The effects of foreign currency translation adjustments are included in stockholders' equity as a component of accumulated other comprehensive income in the accompanying consolidated balance sheets. Transaction gains and losses, which were included in our consolidated statement of operations, amounted to a gain of approximately \$0.8 million, a gain of \$0.4 million, and a loss of \$3.2 million for the fiscal years ended June 30, 2008, 2009, and 2010, respectively.

Earnings per Share—Basic earnings per share is computed by dividing net income available to common shareholders by the weighted average number of common shares outstanding during the period. Diluted earnings per share is computed by dividing net income available to common shareholders by the sum of the weighted average number of common and dilutive potential common shares outstanding. Potential common shares consist of the shares issuable upon the exercise of stock options or warrants under the treasury stock method.

The following table sets forth the computation of basic and diluted earnings per share for the fiscal years ended June 30 (in thousands, except earnings per share data):

	<u>2008</u>	<u>2009</u>	<u>2010</u>
Net income	\$ 13,860	\$ 11,152	\$ 23,557
Effect of dilutive interest in subsidiary stock	—	—	—
Income available to common shareholders	<u>\$ 13,860</u>	<u>\$ 11,152</u>	<u>\$ 23,557</u>
Weighted average shares outstanding—basic	17,428	17,518	17,874
Dilutive effect of stock options and warrants	307	78	515
Weighted average of shares outstanding—diluted	<u>17,735</u>	<u>17,596</u>	<u>18,389</u>
Basic earnings per share	<u>\$ 0.80</u>	<u>\$ 0.64</u>	<u>\$ 1.32</u>
Diluted earnings per share	<u>\$ 0.78</u>	<u>\$ 0.63</u>	<u>\$ 1.28</u>

As of June 30, 2008, 2009 and 2010, approximately 0.4 million, 1.9 million and 0.4 million, respectively, of potentially dilutive shares associated with stock options and stock warrants, collectively, were not included in diluted earnings per common share calculations because to do so would have been antidilutive.

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Provision for Warranties—The Company offers its customers warranties on most products sold to them. These warranties typically provide for repairs and maintenance for a specified time period. Concurrent with the sale of products, a provision for estimated warranty expenses is recorded with a corresponding increase in cost of goods sold. This provision is adjusted periodically based on historical and anticipated experience. Actual expenses of repairs under warranty, including parts and labor, are charged to this provision when incurred.

	Provision for Warranties (in thousands)
Balance on June 30, 2007	7,443
Additions	7,709
Reductions for warranty repair costs	<u>(3,555)</u>
Balance on June 30, 2008	11,597
Additions	4,472
Reductions for warranty repair costs	<u>(5,963)</u>
Balance on June 30, 2009	\$ 10,106
Additions	6,653
Reductions for warranty repair costs	<u>(5,829)</u>
Balance on June 30, 2010	<u>\$ 10,930</u>

Recent Accounting Updates Not Yet Adopted—In October 2009, the Financial Accounting Standards Board issued an accounting standards update amending revenue recognition requirements for multiple-deliverable revenue arrangements. This update provides guidance on separating the deliverables and on the method to measure and allocate arrangement consideration, particularly when the arrangement includes both products and services provided to the customers. The update is effective for revenue arrangements entered into or materially modified in fiscal years beginning on or after June 15, 2010. Early adoption is permitted. The Company has not yet adopted this update and is currently evaluating the impact it may have on its financial condition and results of operations.

In April 2010, the Financial Accounting Standards Board (FASB) issued an accounting standards update relating to milestone revenue recognition. This update provides guidance on the criteria that should be met for determining whether the milestone method of revenue recognition is appropriate primarily as it relates to research and development. The update is effective on a prospective basis for milestones achieved in fiscal years, and interim periods within those years, beginning on or after June 15, 2010. Early adoption is permitted. The Company has not yet adopted this update and is currently evaluating the impact it may have on its financial condition and results of operations.

2. BUSINESS COMBINATIONS

The Company made an acquisition during the first quarter of fiscal year 2010 for cash consideration of \$3.2 million plus consideration contingent upon the financial results of the acquired business over the next four years. The fair value of contingent consideration related to this acquisition was estimated to be \$5.8 million and was recorded at the time of the acquisition as a component of other long-term liabilities. During the third quarter of fiscal 2010, the fair value of this contingent liability was reduced to \$5.1 million. The \$0.7 million associated with this reduction was recorded as other income in the Consolidated Statements of Operations. The acquired business was not considered material to the operations of the Company.

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3. ACCOUNTS RECEIVABLE

Accounts receivable consisted of the following (in thousands):

	June 30,	
	2009	2010
Trade receivables	\$116,140	\$137,502
Receivables related to long term contracts—unbilled costs and accrued profit on progress completed	1,209	1,232
Total	<u>\$117,349</u>	<u>\$138,734</u>
Less: allowance for doubtful accounts	(6,896)	(6,006)
Accounts receivable, net	<u>\$110,453</u>	<u>\$132,728</u>

The unbilled costs and accrued profit at June 30, 2010, are expected to be entirely billed and collected during fiscal 2011.

4. INVENTORIES

Net inventory consisted of the following (in thousands):

	June 30,	
	2009	2010
Raw materials	\$ 77,488	\$ 69,421
Work-in-process	24,648	20,847
Finished goods	48,627	35,662
Total	<u>\$ 150,763</u>	<u>\$ 125,930</u>

5. PROPERTY AND EQUIPMENT

Property and equipment consisted of the following (in thousands):

	Estimated Useful Lives	June 30,	
		2009	2010
Land	N/A	\$ 5,426	\$ 5,078
Buildings	20 years	8,927	8,618
Leasehold improvements	2-20 years	12,628	12,549
Equipment and tooling	3-10 years	48,659	62,861
Furniture and fixtures	3-13 years	4,802	4,753
Computer equipment	3-5 years	16,773	17,738
Computer software	3-10 years	11,032	13,859
Total		108,247	125,456
Less accumulated depreciation and amortization		(66,015)	(73,941)
Property and equipment, net		<u>\$ 42,232</u>	<u>\$ 51,515</u>

During fiscal 2008, 2009 and 2010, depreciation expense was approximately \$15.6 million, \$15.9 million and \$14.4 million, respectively. Included in property and equipment are approximately \$1.6 million and \$1.4 million of assets under capital leases as of June 30, 2009 and 2010, respectively, net of accumulated depreciation.

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6. GOODWILL AND INTANGIBLE ASSETS

The changes in the carrying amount of goodwill for fiscal 2009 and 2010 are as follows (in thousands):

	Security Group	Healthcare Group	Optoelectronics and Manufacturing Group	Consolidated
Balance as of June 30, 2008	\$17,692	\$ 35,569	\$ 7,147	\$ 60,408
Goodwill adjusted during the period		929	226	1,155
Foreign currency translation adjustment	(580)	(762)	(26)	(1,368)
Balance as of June 30, 2009	\$17,112	\$ 35,736	\$ 7,347	\$ 60,195
Goodwill acquired during the period	—	—	4,597	4,597
Foreign currency translation adjustment	(546)	(333)	28	(851)
Balance as of June 30, 2010	<u>\$16,566</u>	<u>\$ 35,403</u>	<u>\$ 11,972</u>	<u>\$ 63,941</u>

Intangible assets subject to amortization consisted of the following (in thousands):

	Weighted Average Lives	June 30, 2009			June 30, 2010		
		Gross Carrying Value	Accumulated Amortization	Intangibles Net	Gross Carrying Value	Accumulated Amortization	Intangibles Net
Amortizable assets:							
Software development costs	5 years	\$ 9,754	\$ 3,198	\$ 6,556	\$ 11,877	\$ 3,954	\$ 7,923
Patents	9 years	921	334	587	1,630	388	1,242
Core technology	10 years	2,224	977	1,247	2,029	1,094	935
Developed technology	12 years	17,360	7,169	10,191	17,246	8,942	8,304
Customer relationships/ backlog	7 years	9,456	4,876	4,580	10,437	6,132	4,305
Total amortizable assets		39,715	16,554	23,161	43,219	20,510	22,709
Non-amortizable assets:							
Trademarks		9,290		9,290	9,266		9,266
Total intangible assets		<u>\$ 49,005</u>	<u>\$ 16,554</u>	<u>\$ 32,451</u>	<u>\$ 52,485</u>	<u>\$ 20,510</u>	<u>\$ 31,975</u>

Amortization expense for the fiscal 2008, 2009 and 2010 was \$3.7 million, \$3.9 million and \$4.1 million, respectively. Future acquisitions could cause these amounts to increase. At June 30, 2010, estimated future amortization expense was as follows (in thousands):

2011	\$ 4,091
2012	4,058
2013	3,768
2014	2,584
2015	1,213
2016 and thereafter	6,995
Total	<u>\$22,709</u>

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Software development costs for software products incurred before establishing technological feasibility, are charged to operations. Software development costs incurred after establishing technological feasibility and purchased software costs are capitalized on a product-by-product basis until the product is available for general release to customers at which time amortization begins. Annual amortization, charged to cost of sales, is the greater of: (i) the amount computed using the ratio that current gross revenues for a product bear to the total current and anticipated future gross revenues for that product or (ii) the straight-line method over the remaining estimated economic life of the product. During fiscal 2008, 2009 and 2010, the Company capitalized software development costs in the amount of \$1.9 million, \$3.4 million and \$2.1 million, respectively.

7. RESTRUCTURING AND OTHER CHARGES

In response to challenging worldwide economic conditions, the Company continued to optimize its cost structure by reducing excess workforce and facilities and consolidating and relocating certain manufacturing facilities. Such efforts resulted in restructuring charges of \$4.7 million in 2008, \$7.1 million in 2009 and \$2.9 million in 2010. The following table analyzes the key components of these restructuring and other charges throughout fiscal 2008, 2009 and 2010:

	Security Division	Healthcare Division	Optoelectronics and Manufacturing Division	Corporate	Consolidated
Accrued balance as of June 30, 2007	\$ 654	\$ 201	\$ 171	\$ —	\$ 1,026
Expensed during the year					
Facility closure	890	1,145	78	—	2,113
Employee termination costs	1,413	813	349	—	2,575
Total expensed during year	2,303	1,958	427	—	4,688
Paid during the year	2,633	1,340	577	—	4,550
Accrued balance as of June 30, 2008	<u>\$ 324</u>	<u>\$ 819</u>	<u>\$ 21</u>	<u>\$ —</u>	<u>\$ 1,164</u>
Expensed during the year					
Facility closure	577	1,502	166	—	2,245
Employee termination costs	673	1,829	76	300	2,878
Litigation	—	—	—	2,000	2,000
Total expensed during year	1,250	3,331	242	2,300	7,123
Paid during the year	876	4,072	64	140	5,152
Accrued balance as of June 30, 2009	<u>\$ 698</u>	<u>\$ 78</u>	<u>\$ 199</u>	<u>\$ 2,160</u>	<u>\$ 3,135</u>
Expensed during the year					
Facility closure	509	89	559	—	1,157
Employee termination costs	6	1,210	396	90	1,702
Total expensed during year	515	1,299	955	90	2,859
Paid during the year	750	644	854	2,250	4,498
Accrued balance as of June 30, 2010	<u>\$ 463</u>	<u>\$ 733</u>	<u>\$ 300</u>	<u>\$ —</u>	<u>\$ 1,496</u>

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8. LINE-OF-CREDIT BORROWINGS AND DEBT

The Company maintains a credit agreement with certain lenders allowing for initial borrowings of up to \$124.5 million. The credit agreement consists of a \$74.5 million, five-year, revolving credit facility (including a \$64.5 million sub-limit for letters-of-credit) and a \$50 million five-year term loan. Borrowings under the agreement bear interest at either (i) the London Interbank Offered Rate (LIBOR) plus between 2.00% and 2.50% or (ii) the bank's prime rate plus between 1.00% and 1.50%. The rates are determined based on the Company's consolidated leverage ratio. As of June 30, 2010, the weighted-average interest rate under the credit agreement was 2.35%. The Company's borrowings under the credit agreement are guaranteed by the Company's domestic subsidiaries and are secured by substantially all of the Company's and its subsidiary guarantors' assets. The agreement contains various representations, warranties, affirmative, negative and financial covenants, and conditions of default customary for financing agreements of this type, including restrictions on the Company's ability to pay cash dividends. As of June 30, 2010, \$32.3 million was outstanding under the term loan, while no debt was outstanding under the revolving credit facility, and \$38.4 million was outstanding under the letter-of-credit facility.

Several of the Company's foreign subsidiaries maintain bank lines-of-credit, denominated in local currencies, to meet short-term working capital requirements and for the issuance of letters-of-credit. As of June 30, 2010, \$10.0 million was outstanding under these letter-of-credit facilities, while no debt was outstanding. As of June 30, 2010, the total amount available under these credit facilities was \$18.0 million, with a total cash borrowing sub-limit of \$4.0 million.

In fiscal 2005, the Company entered into a bank loan of \$5.3 million to fund the acquisition of land and buildings in the U.K. The loan is payable over a 20-year period. The loan bears interest at British pound-based LIBOR plus 1.2%, payable on a quarterly basis. As of June 30, 2010, \$3.0 million remained outstanding under this loan at an interest rate of 1.9% per annum.

Long-term debt consisted of the following at June 30 (in thousands):

	<u>2009</u>	<u>2010</u>
Five-year term loan due in 2013	\$ 42,763	\$ 32,281
Twenty-year term loan due in 2024	3,533	3,015
Capital leases	1,354	711
Other	710	102
	<u>48,360</u>	<u>36,109</u>
Less current portion of long-term debt	8,557	12,743
Long-term portion of debt	<u>\$ 39,803</u>	<u>\$ 23,366</u>

Fiscal year principal payments of long-term debt as of June 30, 2010 are as follows (in thousands):

2011	\$ 12,743
2012	6,294
2013	14,609
2014	218
2015	218
2016 and thereafter	2,027
Total	<u>\$ 36,109</u>

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9. STOCK-BASED COMPENSATION

As of June 30, 2010, the Company maintained one significant stock-based compensation plan – the 2006 Equity Participation Plan of OSI Systems (OSI Plan). The OSI Plan allows for the issuance of restricted stock and the granting of stock options. As of June 30, 2007, the Company had maintained three significant plans: (a) the OSI Plan, (b) the 2005 Equity Participation Plan of Spacelabs Healthcare (Spacelabs Plan) and (c) the 2006 Equity Participation Plan of Rapiscan Systems Holdings, Inc. (Rapiscan Plan). However, during fiscal 2008, the Company converted all of the options outstanding under the Spacelabs Plan and Rapiscan Plan into options under the OSI Plan. The methodology used for such conversions provided equivalent fair values under the OSI Plan. Therefore, no additional compensation expense was incurred by the Company as a result of these conversions.

The Company recorded stock-based-compensation expense in the consolidated statement of operations as follows (in thousands):

	<u>2008</u>	<u>2009</u>	<u>2010</u>
Cost of goods sold	\$ 215	\$ 291	\$ 287
Selling, general and administrative	4,331	4,527	4,499
Research and development	231	237	225
Stock based compensation expense before taxes	4,777	5,055	5,011
Related income tax benefit	1,504	1,819	1,750
Stock based compensation expense, net of estimated taxes	<u>\$3,273</u>	<u>\$3,236</u>	<u>\$3,261</u>

As of June 30, 2010, total unrecognized compensation cost related to non-vested stock-based compensation grants amounted to \$1.6 million for stock options and \$5.5 million for restricted stock under the OSI Plan. The Company expects to recognize these costs over a weighted-average period of 1.8 years with respect to the options and 2.7 years for grants of restricted stock.

Employee Stock Purchase Plan—The Company has an employee stock purchase plan under which eligible employees may purchase a limited number of shares of Common Stock at a discount of up to 15% of the market value of such stock at pre-determined, plan-defined dates. During the three years ended June 30, 2008, 2009 and 2010, employees purchased 65,908, 75,594 and 141,136 shares, respectively. As of June 30, 2010, there were 1,295,063 shares of the Company's Common Stock available for issuance under the plan.

OSI Plan

Stock Options—Under the OSI Plan, the Company is authorized to grant up to 5,350,000 shares of Common Stock in the form of incentive options, nonqualified options or restricted stock. Under the plan, the exercise price of nonqualified options may not be less than 85% of the fair market value of the Company's Common Stock on the date of grant. The exercise price of incentive stock options may not be less than the fair market value of the Company's Common Stock at the date of grant. The exercise price of incentive stock options granted to individuals who own more than 10% of the Company's voting stock may not be less than 110% of the fair market value of the Company's Common Stock on the date of grant.

No single method of estimating volatility is proper under all circumstances and that to the extent a company can derive implied volatility based on the trading of its financial instruments on a public market, it may be appropriate to use both implied and historical volatility in its assumptions. The Company has certain financial

OSI SYSTEMS, INC. AND SUBSIDIARIES
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instruments that are publicly traded from which the Company can derive the implied volatility. Therefore, the Company used implied and historical volatility for valuing its stock options. The Company believes that implied and historical volatility is a better indicator of expected volatility because it is generally reflective of both historical volatility and expectations of how future volatility will differ from historical volatility.

The Company determined the fair value of options issued during fiscal 2008, 2009 and 2010 as of the date of the grant, using the Black-Scholes option pricing model with the following weighted average assumptions:

	<u>2008</u>	<u>2009</u>	<u>2010</u>
Expected dividend	0%	0%	0%
Risk-free interest rate	4.1%	1.8%	2.0%
Expected volatility	40.5%	41.5%	39.0%
Expected life (in years)	4.1	4.3	4.3

The following summarizes stock option activity for fiscal years 2008, 2009 and 2010:

	<u>Number of Options</u>	<u>Weighted- Average Exercise Price</u>	<u>Weighted-Average Remaining Contractual Term</u>	<u>Aggregate Intrinsic Value (\$000)</u>
Outstanding at June 30, 2007	1,332,129	18.63		
Granted	279,000	20.95		
Converted from Spacelabs Plan	456,226	19.69		
Converted from Rapiscan Plan	622,309	17.06		
Exercised	(340,642)	16.62		
Expired or cancelled	(58,026)	21.20		
Outstanding at June 30, 2008	2,290,996	18.93		
Granted	332,500	14.32		
Exercised	(163,680)	15.21		
Expired or cancelled	(283,953)	19.51		
Outstanding at June 30, 2009	2,175,863	17.69		
Granted	257,500	16.41		
Exercised	(660,764)	16.98		
Expired or cancelled	(216,489)	20.01		
Outstanding at June 30, 2010	<u>1,556,110</u>	<u>17.46</u>	<u>4.9 years</u>	<u>\$ 16,087</u>
Exercisable at June 30, 2010	<u>1,053,569</u>	<u>\$ 18.32</u>	<u>3.1 years</u>	<u>\$ 9,992</u>

The per-share weighted-average grant-date fair value of stock options granted under the OSI Plan was \$7.74, \$5.16 and \$5.64 for fiscal 2008, 2009 and 2010, respectively. The total intrinsic value of options exercised during fiscal 2010 was \$5.9 million.

In fiscal 2008, the Company converted 5,900,385 options under the Spacelabs Plan into 456,226 options under the OSI Plan and converted 7,221,000 options under the Rapiscan Plan into 622,309 options under the OSI Plan. In both of these cases, no additional compensation expense was required to be recorded.

Restricted Stock Awards—Under the OSI Plan, the Company granted 202,700, 227,626 and 247,800 restricted shares during fiscal 2008, 2009 and 2010, respectively.

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A summary of restricted stock award activity for the periods indicated was as follows:

	Shares	Weighted-Average Fair Value
Nonvested at June 30, 2007	—	\$ —
Granted	202,700	23.01
Vested	(14,982)	22.59
Forfeited	(250)	23.18
Nonvested at June 30, 2008	187,468	\$ 23.04
Granted	227,626	13.56
Vested	(43,695)	23.35
Forfeited	(11,708)	20.97
Nonvested at June 30, 2009	359,691	\$ 17.07
Granted	247,800	18.20
Vested	(112,665)	17.99
Forfeited	(15,289)	17.18
Nonvested at June 30, 2010	<u>479,537</u>	<u>\$ 17.44</u>

The per-share weighted average grant-date fair value of restricted stock granted under the OSI Plan was \$23.01, \$13.56 and \$18.20 for fiscal 2008 and 2009 and 2010, respectively. The total fair value of shares vested during fiscal 2008, 2009 and 2010 was \$0.3 million, \$1.0 million and \$2.0 million, respectively.

As of June 30, 2010, there were 524,017 shares available for grant under the OSI Plan. Under the terms of the OSI Plan, no more than 349,121 of these shares may be granted in the form of restricted stock.

10. INCOME TAXES

The following is a geographical breakdown of income (loss) before the provision (benefit) for income taxes (in thousands):

	2008	2009	2010
Pre-tax income (loss):			
United States	\$ 665	\$ (4,751)	\$14,888
Foreign	13,774	21,296	20,108
Total pre-tax income	<u>\$14,439</u>	<u>\$16,545</u>	<u>\$34,996</u>

The Company's provision for income taxes consists of the following (in thousands):

	2008	2009	2010
Current:			
Federal	\$ 2,433	\$ 3,442	\$ 2,809
State	(794)	418	745
Foreign	2,190	3,556	2,644
Total current provision	<u>3,829</u>	<u>7,416</u>	<u>6,198</u>
Deferred:			
Federal	\$(5,237)	\$(2,384)	\$ 2,640
State	1,281	(121)	(374)
Foreign	706	482	2,975
Total deferred provision	<u>(3,250)</u>	<u>(2,023)</u>	<u>5,241</u>
Total provision	<u>\$ 579</u>	<u>\$ 5,393</u>	<u>\$11,439</u>

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The exercise of stock options during the fiscal year ended June 30, 2008, 2009 and 2010 resulted in additional tax benefit or provision and has been reflected as an adjustment of income taxes payable and an adjustment of additional paid-in capital. The adjustments recorded were \$(0.4) million, \$(0.6) million and \$0.7 million for the years ended June 30, 2008, 2009 and 2010, respectively.

On July 1, 2007, the Company adopted the accounting standards update related to accounting for uncertainty in income taxes. The cumulative effect of applying this accounting standards update to the Company was recorded as a decrease of \$3.3 million to retained earnings, an increase of \$2.4 million to deferred tax asset and an increase of \$6.2 million to deferred tax liability.

As of June 30, 2008, June 30, 2009 and June 30, 2010, the Company's liability for uncertain tax positions was \$6.5 million, \$8.4 million and \$9.4 million, respectively. Of the \$9.4 million, \$7.0 million represents the amount of unrecognized tax benefits that, if recognized, would affect the effective tax rate.

The Company recognizes potential interest and penalties related to income tax matters in income tax expense. As of June 30, 2010, the Company has \$2.5 million accrued for the payment of interest and penalties. The Company's uncertain tax positions are related to tax years that remain subject to examination by the relevant tax authorities. These include the fiscal years after 2006 for federal purposes, the fiscal years after 2005 for state purposes and fiscal years after 1998 for various foreign jurisdictions. Facts and circumstances could arise in the twelve month period following June 30, 2010 that could cause the Company to reduce the liability for unrecognized tax benefits, including but not limited to, settlement of income tax positions or expiration of statute of limitations. Since the ultimate resolution of uncertain tax positions depends on many factors and assumptions, the Company is not able to estimate the range of potential changes in the liability for unrecognized tax benefits or the timing of such changes.

A summary of activity of unrecognized tax benefits for fiscal 2008, 2009 and 2010 was as follows (in thousands).

Balance as July 1, 2007	\$4,373
Additions on tax positions for the current year	463
Reductions as a result of a lapse of statute of limitations	(69)
Balance as June 30, 2008	\$4,767
Additions on tax positions for the current year	148
Additions on tax positions from prior years	972
Balance at June 30, 2009	\$5,887
Additions on tax positions for the current year	1,119
Additions on tax positions from prior years	780
Reductions as a result of a lapse of statute of limitations	(456)
Reduction in tax position from prior year	(453)
Balance at June 30, 2010	<u>\$6,877</u>

The Company does not provide for U.S. income taxes on the undistributed earnings of its foreign subsidiaries as it is the Company's intention to utilize those earnings in the foreign operations for an indefinite period of time. At June 30, 2010, undistributed earnings of the foreign subsidiaries amounted to approximately \$91.5 million. It is not practical to determine the amount of income or withholding tax that would be payable upon the remittance of these earnings.

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Included within the tax benefit for fiscal 2008 is a net tax benefit of \$4.0 million as a result of discrete items impacting the tax provision, the largest of which was a \$4.3 million tax benefit associated with the repurchase of the minority interest of Spacelabs Healthcare (see Note 1), which the Company completed during the second quarter of fiscal 2008.

Deferred income tax assets (liabilities) consisted of the following (in thousands):

	June 30,	
	2009	2010
Deferred income tax assets:		
R&D tax credit carryforwards	\$ 1,629	\$ 3,830
Net operating loss carryforwards	3,915	3,475
Revitalization zone deductions	879	879
Allowance for doubtful accounts	1,585	1,206
Inventory reserve	7,736	8,014
Inventory capitalization	3,615	3,615
Accrued liabilities	4,856	4,772
Deferred compensation	5,353	4,839
Unrecognized tax benefit	2,435	2,435
Other assets	8,892	6,336
Total deferred income tax assets	40,895	39,401
Valuation allowance	(6,223)	(6,352)
Net deferred income tax assets	34,672	33,049
Deferred income tax liabilities:		
Depreciation	(378)	(314)
State income taxes	(1,497)	(1,305)
Amortization of intangible assets	(8,695)	(10,554)
Other liabilities	(155)	(30)
Total deferred income tax liabilities	(10,725)	(12,203)
Net deferred tax asset	<u>\$ 23,947</u>	<u>\$ 20,846</u>

As of June 30, 2010, the Company had federal net operating loss carry forwards of approximately \$4.1 million and state net operating loss carry forwards of \$1.1 million. The Company's federal net operating losses will begin to expire in the tax year ending June 30, 2018, and are subject to limitations on their utilization. In addition, the Company had state tax credit carry forwards, including research and development and revitalization zone credits, of approximately \$2.5 million. The Company's state tax credit carry forwards will begin to expire in the tax year ending June 30, 2011. As of June 30, 2010, the Company has a federal research and development tax credit carry forward of approximately \$2.2 million. The Company's federal credit carry forwards will begin to expire in the tax year ending June 30, 2019.

The Company has established a valuation allowance in accordance with the provisions outlined in the accounting standards related to income taxes. The valuation allowance relates to the net operating loss of foreign subsidiaries, net operating loss subject to Separate Return Limitation Year rules, an unrealized capital loss related to a write-down of an equity investment and revitalization zone credits. During the year ended June 30, 2010, the

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Company recorded a \$0.1 million increase to its valuation allowance, which entirely related to foreign NOL. The Company reviews the adequacy of valuation allowances and releases the allowances when it is determined that it is more likely than not that the benefits will be realized.

The consolidated effective income tax rate differs from the federal statutory income tax rate due primarily to the following:

	June 30,		
	2008	2009	2010
Provision (benefit) for income taxes at federal statutory rate	(35.0)%	35.0%	35.0%
State income taxes and credits—net of federal benefit	2.2	1.2	0.7
Research and development tax credits	(2.0)	(2.8)	(3.2)
Foreign tax credits	(1.0)	(3.4)	(0.6)
Subpart F income	5.8	4.1	0.6
Stock options adjustment	1.4	0.8	(0.5)
Foreign income subject to tax at other than federal statutory rate	(25.8)	(18.2)	(5.3)
Nondeductible expenses	2.7	1.1	0.1
Change in valuation allowance	12.3	2.6	1.0
Reversal of Spacelabs deferred tax liability	(29.6)	—	—
Unrecognized tax benefit	2.3	12.2	1.8
Other	0.7	—	3.1
Effective income tax rate	<u>4.0%</u>	<u>32.6%</u>	<u>32.7%</u>

The provision for income taxes consists of provisions for federal, state, and foreign income taxes. The Company operates in an international environment with significant operations in various locations outside the U.S. Accordingly, the consolidated income tax rate is a composite rate reflecting the earnings in the various locations and the applicable rates.

11. COMMITMENTS AND CONTINGENCIES

The following is a summary of commitments as of June 30, 2010 (in thousands):

	Payments Due by Period				
	Total	Less than 1 year	2-3 years	4- 5 years	After 5 years
Total debt (excluding capital lease obligations)	\$35,398	\$11,920	\$21,015	\$ 436	\$2,027
Capital lease obligations	\$ 711	\$ 711	\$ —	\$ —	\$ —
Operating leases	\$38,197	\$10,794	\$17,197	\$9,749	\$ 457
Defined benefit plan obligation	\$ 7,921	\$ 879	\$ 550	\$ 514	\$5,978

Operating Leases—The Company leases facilities and certain equipment under various operating lease agreements. Certain leases provide for periodic rent increases and may contain escalation clauses and renewal options. Rent expense totaled \$10.0 million, \$9.6 million and \$10.4 million for fiscal years 2008, 2009 and 2010, respectively.

Commitments—Under the terms and conditions of the purchase agreements associated with the following acquisitions, the Company may be obligated to make additional payments.

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In fiscal 2003, the Company purchased a minority equity interest in CXR Limited. In June 2004, the Company increased its equity interest to approximately 75% and in December 2004, the Company acquired the remaining 25%. As compensation to the selling shareholders for this remaining interest, the Company agreed to make certain royalty payments during the 18 years following the acquisition of this remaining interest. Royalty payments are based on the license of, or sales of products containing, technology owned by CXR Limited. As of June 30, 2010, no royalty payments have been earned.

In fiscal 2004, the Company acquired Advanced Research & Applications Corp. During the seven years following the acquisition, contingent consideration is payable based on its net revenues, provided certain requirements are met. The contingent consideration is capped at \$30.0 million. As of June 30, 2010, no contingent consideration has been earned.

In fiscal 2006, the Company acquired certain assets of InnerStep, B.S.E., Inc. During the seven years following the acquisition, contingent consideration is payable based on its profits before interest and taxes, provided certain requirements are met. The contingent consideration is capped at \$6.0 million. As of June 30, 2010, no contingent consideration has been earned.

In fiscal 2009, the Company acquired a company that offers services in connection with security inspection products. Contingent consideration is payable based on net receipts generated from new business during the three years following the acquisition, provided certain requirements are met. The contingent consideration is capped at \$10.0 million. As of June 30, 2010, no contingent consideration has been earned.

In fiscal 2010, the Company completed an acquisition in which the fair value of contingent consideration related to this acquisition was estimated to be \$5.8 million and was recorded at the time of the acquisition as a component of other long-term liabilities. During the third quarter of fiscal 2010, the fair value of this contingent liability was reduced to \$5.1 million. The \$0.7 million associated with this reduction was recorded as other income in the Consolidated Statements of Operations.

Environmental Contingencies—The Company is subject to various environmental laws. The Company's practice is to ensure that Phase I environmental site assessments are conducted for each of its properties in the United States at which the Company manufactures products in order to identify, as of the date of such report, potential areas of environmental concern related to past and present activities or from nearby operations. In certain cases, the Company has conducted further environmental assessments consisting of soil and groundwater testing and other investigations deemed appropriate by independent environmental consultants.

During one investigation, the Company discovered soil and groundwater contamination at its Hawthorne, California facility. The Company filed the requisite reports concerning this problem with the appropriate environmental authorities in fiscal 2001. The Company has not yet received any response to such reports, and no agency action or litigation is presently pending or threatened. The Company's site was previously used by other companies for semiconductor manufacturing similar to that presently conducted on the site by us, and it is not presently known who is responsible for the contamination or, if required, the remediation. The groundwater contamination is a known regional problem, not limited to the Company's premises or its immediate surroundings.

The Company has also been informed of soil and groundwater evaluation efforts at a facility that its Ferson Technologies subsidiary previously leased in Ocean Springs, Mississippi. Ferson Technologies occupied the facility until October 2003. The Company believes that the owner and previous occupants of the facility have primary responsibility for any remediation that may be required and have an agreement with the facility's owner

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under which the owner is responsible for remediation of pre-existing conditions. However, as site evaluation efforts are still in progress, and may be for some time, the Company is unable at this time to ascertain whether Ferson Technologies bears any exposure for remediation costs under applicable environmental regulations.

The Company has not accrued for loss contingencies relating to the above environmental matters because it believes that, although unfavorable outcomes may be possible, they are not considered by the Company's management to be probable and reasonably estimable.

If one or more of these matters are resolved in a manner adverse to the Company, the impact on the Company's results of operations, financial position and/or liquidity could be material.

Legal Proceedings—The Company is involved in various claims and legal proceedings arising out of the ordinary course of business. In the Company's opinion after consultation with legal counsel, the ultimate disposition of such proceedings is not likely to have a material adverse effect on its financial position, future results of operations, or cash flows. The Company has not accrued for loss contingencies relating to such matters because the Company believes that, although unfavorable outcomes in the proceedings may be possible, they are not considered by management to be probable or reasonably estimable. If one or more of these matters are resolved in a manner adverse to the Company, the impact on the Company's results of operations, financial position and/or liquidity could be material.

12. SHAREHOLDERS' EQUITY

Stock Repurchase Program

The Company's Board of Directors has authorized a Common Stock repurchase program. During fiscal 2009, the Company repurchased 619,768 shares under this program. No shares were repurchased under this program in fiscal 2008 or 2010. At June 30, 2010, 711,205 shares were available for repurchase under the stock repurchase program. There is no timeframe to complete the repurchase program. Upon repurchase, shares were restored to the status of authorized but unissued shares in the accompanying Consolidated Financial Statements.

Warrants

In June 2004, the Company issued and sold an aggregate of 1,500,000 shares of Common Stock in a private placement to institutional investors and received net proceeds of \$31 million. As part of the transaction, the Company issued warrants to purchase 337,500 additional shares of the Company's Common Stock at an exercise price of \$27.73 per share exercisable at any time, in full or part, expiring on June 1, 2011.

13. Related-Party Transactions

In 1994, the Company, together with an unrelated company, formed ECIL-Rapiscan Security Products Limited, a joint venture organized under the laws of India. The Company owns a 36% interest in the joint venture, the Company's Chairman and Chief Executive Officer owns a 10.5% interest, and the Company's Executive Vice President and President of the Company's Security division owns a 4.5% ownership interest. The Company's initial investment was approximately \$0.1 million. For the years ended June 30, 2008, 2009 and 2010, the Company's equity earnings in the joint venture were approximately to \$0.4 million, \$0.5 million and \$0.4 million, respectively. The Company, its Chairman and Chief Executive Officer and the Company's Executive Vice President and President of the Company's Security division collectively control less than 50% of the board of directors voting power in the joint venture. As a result, the Company accounts for the investment

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under the equity method of accounting. The joint venture was formed for the purpose of the manufacture, assembly, service and testing of security and inspection systems and other products. Some of the Company's subsidiaries are suppliers to the joint venture partner, which in turn manufactures and sells the resulting products. Sales to the joint venture partner for fiscal 2008, 2009 and 2010 were approximately \$1.6 million, \$4.4 million and \$4.4 million, respectively. Receivables from the joint venture were \$2.7 million and \$0.9 million as of June 30, 2009 and 2010, respectively.

The Company has contracted with entities owned by members of its Board of Directors and/or their families to provide messenger service, auto rental and printing services. Such expenses for 2008, 2009 and 2010 were approximately \$54,000, \$54,000 and \$64,000 for messenger services and auto rental and \$42,000, \$45,000 and \$60,000 for printing services, respectively.

14. Employee Benefit Plans

Employee Retirement Savings Plans

The Company has various qualified employee retirement savings plans. Participants can contribute certain amounts to the plans and the Company matches a certain portion of employee contributions. The Company contributed approximately \$2.3 million, \$2.8 million and \$3.1 million to the plans for the fiscal years ended June 30, 2008, 2009 and 2010, respectively.

Deferred Compensation Plan

In May 2008, the Company adopted a deferred compensation plan, which met the requirements for deferred compensation under Section 409A of the Internal Revenue Code. The plan provides that selected employees are eligible to defer up to 80% of their salaries and up to 100% of their bonuses. The Company may also make employer contributions to participant accounts in certain circumstances. The benefits under this plan are unsecured. Participants are generally eligible to receive payment of their vested benefit at the end of their elected deferral period or after termination of their employment for any reason or at a later date to comply with the restrictions of Section 409A. Discretionary company contributions and the related earnings are subject to a vesting schedule dependent upon years of service to the Company and, also, vest completely upon the participant's disability, death or a change of control. The Company made no employer contributions to the plan during fiscal year 2008 and made contributions of \$0.6 million and \$0.5 million during fiscal year 2009 and 2010, respectively. As of June 30, 2010, the Company held assets of \$3.7 million and liabilities of \$3.5 million. Assets related to this plan are included in Other Assets and liabilities related to this plan are included in Other Long Term Liabilities on the balance sheets. The plan liabilities include accrued employer contributions not yet funded to the plan.

Employee Pension Plans

The Company sponsors a number of qualified and nonqualified pension plans for its employees at certain locations. In accordance with accounting standards for employee pension and postretirement benefits, the Company fully recognizes the overfunded or underfunded status of each of its defined benefit plan as an asset or liability in the consolidated balance sheet. The asset or liability equals the difference between the fair value of the plan's assets and its benefit obligation. The liabilities associated with underfunded plans are classified as noncurrent, except to the extent the fair value of the plan's assets is less than the plan's estimated benefit payments over the next 12 months. The Company measures its pension and postretirement benefit plan assets and benefit obligations as of June 30.

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During fiscal 2009, the Company recognized curtailment gains of \$0.3 million due to headcount reductions in foreign locations related to participants in defined benefit plans.

The following provides a reconciliation of the changes in the plans' benefit obligations and fair value of assets for fiscal years 2009 and 2010, and a statement of the funded status as of June 30, 2009 and 2010 (in thousands):

	<u>2009</u>	<u>2010</u>
Change in Benefit Obligation		
Benefit obligation at beginning of year	\$ 8,977	\$ 7,997
Translation adjustment	(1,939)	(459)
Service costs	2,093	68
Interest costs	537	502
Curtailment / plan amendment	(333)	460
Plan participants' contributions	(88)	(51)
Actuarial loss (gain)	(1,198)	901
Benefits paid	(52)	(53)
Benefit obligation at end of year	<u>7,997</u>	<u>9,365</u>
Change in Plan Assets		
Fair value of plan assets at beginning of year	5,829	4,316
Translation adjustment	(1,108)	47
Actual return on plan assets	(814)	651
Company contributions	528	299
Plan participants' contributions	—	65
Benefits paid	(119)	(124)
Fair value of plan assets at end of year	<u>4,316</u>	<u>5,254</u>
Funded status	<u>(3,681)</u>	<u>(4,111)</u>
Unrecognized net actuarial loss	—	—
Net amount recognized	<u><u>\$(3,681)</u></u>	<u><u>\$(4,111)</u></u>
Amount recognized in Consolidated Balance Sheets consist of:		
Investments	\$ 535	\$ 552
Accrued pension liability	(4,146)	(4,883)
Accumulated other comprehensive income	3,088	3,072

The following table provides the net periodic benefit costs for each of the fiscal years ended June 30, (in thousands):

	<u>2008</u>	<u>2009</u>	<u>2010</u>
Net Periodic Benefit Costs			
Service costs	\$ 259	\$ 297	\$ 49
Interest costs	486	537	464
Expected return on plan assets	(220)	(300)	(306)
Amortization of prior service costs	—	148	258
Recognized actuarial loss/(gain)	90	(135)	284
Curtailment	—	(333)	(14)
Net periodic benefit cost	<u><u>\$ 615</u></u>	<u><u>\$ 214</u></u>	<u><u>\$ 735</u></u>

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Plan Assumptions

	<u>2009</u>	<u>2010</u>
Weighted average assumptions at year-end:		
Discount rate	6.0%	5.1%
Expected return on plan assets	6.6%	6.2%
Rate of compensation increase	0.5%	0.2%

The long term return on assets has been derived from the weighted average of assumed returns on each of the major asset categories. The weighted average is based on the actual proportion of each major asset class held, rather than a benchmark portfolio of assets. The expected returns for each major asset class have been derived from a combination of both historical market returns and current market data as well as the views of a range of investment managers.

Plan Assets and Investment Policy

	<u>Fiscal year ended</u> <u>June 30, 2009</u>		<u>Fiscal year ended</u> <u>June 30, 2010</u>	
	<u>Proportion of</u> <u>Fair Value</u>	<u>Expected Rate</u> <u>of Return</u>	<u>Proportion of</u> <u>Fair Value</u>	<u>Expected Rate</u> <u>of Return</u>
Equity securities	71%	7%	83%	7%
Debt securities	26%	6%	16%	3%
Other	3%	5%	1%	5%
Combined	<u>100%</u>	6.6%	<u>100%</u>	6.2%

The defined benefit plans' assets are invested in a range of pooled investment funds that provide access to a diverse range of asset classes. The investment objective is to maximize the investment return over the long term without exposing the fund to an unnecessary level of risk. Within this objective, it is recognized that benefits will be secured by the purchase of annuities at the time of employee retirement.

The benchmark is to hold assets broadly in the proportion of 50% equity securities and 50% debt securities. This proportion is allowed to fluctuate with market movements and is not formally rebalanced. The equity holdings are maintained in balanced funds under the control of investment managers.

Day-to-day equities selection decisions are delegated to investment managers, although these are monitored against performance and risk targets. Due to the nature of the pooled funds, there are no significant holdings in any single company (greater than 5% of the total assets). The investment strategy is reviewed on a regular basis, based on the results of the liability studies.

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Projected Benefit Payments

The following table reflects estimated benefits payments, based upon the same assumptions used to measure the benefit obligation and net pension cost, as of June 30, 2010 (in thousands):

	<u>Pension Benefits</u>
July 1, 2010 to June 30, 2011	879
July 1, 2011 to June 30, 2012	375
July 1, 2012 to June 30, 2013	175
July 1, 2013 to June 30, 2014	333
July 1, 2014 to June 30, 2015	181
July 1, 2015 to June 30, 2020	5,978

Company Contribution

Currently, the Company's weighted average contribution rate is 2% of pensionable salaries. If the Company contributions continue at the current rate, the estimated total Company contributions for fiscal 2011 will be approximately \$0.3 million.

15. SEGMENT INFORMATION

The Company has determined that it operates in three identifiable industry segments, (a) security and inspection systems (Security division), (b) medical monitoring and anesthesia systems (Healthcare division), and (c) optoelectronic devices and manufacturing (Optoelectronics and Manufacturing division). The Company also has a corporate segment (Corporate) that includes executive compensation and certain other general and administrative expenses, expenses related to stock issuances and legal and audit and other professional service fees not allocated to product segments. Both the Security and Healthcare divisions comprise primarily end-product businesses whereas the businesses of the Optoelectronics and Manufacturing division primarily supply components and subsystems to original equipment manufacturers, including to the Security and Healthcare divisions. Sales between divisions are at transfer prices that approximate market values. All other accounting policies of the segments are the same as described in Note 1, Summary of Significant Accounting Policies.

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The following tables present the operations and identifiable assets by industry segment (in thousands):

	2008					
	Security Division	Healthcare Division	Optoelectronics and Manufacturing Division	Corporate	Eliminations	Consolidated
Revenues:						
External customer revenue	\$ 225,836	\$ 256,695	\$ 140,557	\$ —	\$ —	\$ 623,088
Revenue between product segments	—	—	47,067	—	(47,067)	—
Total revenues	<u>\$ 225,836</u>	<u>\$ 256,695</u>	<u>\$ 187,624</u>	<u>\$ —</u>	<u>\$ (47,067)</u>	<u>\$ 623,088</u>
Income (loss) from operations	<u>\$ 5,365</u>	<u>\$ 12,918</u>	<u>\$ 13,114</u>	<u>\$ (12,290)</u>	<u>\$ (199)</u>	<u>\$ 18,908</u>
Segments assets	<u>\$ 199,884</u>	<u>\$ 172,038</u>	<u>\$ 95,615</u>	<u>\$ 43,313</u>	<u>\$ (3,209)</u>	<u>\$ 507,641</u>
Capital expenditures	<u>\$ 3,939</u>	<u>\$ 5,080</u>	<u>\$ 2,149</u>	<u>\$ 949</u>	<u>\$ —</u>	<u>\$ 12,117</u>
Depreciation and amortization	<u>\$ 6,233</u>	<u>\$ 9,004</u>	<u>\$ 3,580</u>	<u>\$ 525</u>	<u>\$ —</u>	<u>\$ 19,342</u>
	2009					
	Security Division	Healthcare Division	Optoelectronics and Manufacturing Division	Corporate	Eliminations	Consolidated
Revenues:						
External customer revenue	\$ 240,919	\$ 214,260	\$ 135,182	\$ —	\$ —	\$ 590,361
Revenue between product segments	—	—	45,941	—	(45,941)	—
Total revenues	<u>\$ 240,919</u>	<u>\$ 214,260</u>	<u>\$ 181,123</u>	<u>\$ —</u>	<u>\$ (45,941)</u>	<u>\$ 590,361</u>
Income (loss) from operations	<u>\$ 14,324</u>	<u>\$ 5,106</u>	<u>\$ 14,501</u>	<u>\$ (13,890)</u>	<u>\$ (560)</u>	<u>\$ 19,481</u>
Segments assets	<u>\$ 191,164</u>	<u>\$ 155,366</u>	<u>\$ 84,434</u>	<u>\$ 47,633</u>	<u>\$ (3,769)</u>	<u>\$ 474,828</u>
Capital expenditures	<u>\$ 2,322</u>	<u>\$ 2,890</u>	<u>\$ 4,293</u>	<u>\$ 1,347</u>	<u>\$ —</u>	<u>\$ 10,852</u>
Depreciation and amortization	<u>\$ 5,774</u>	<u>\$ 8,389</u>	<u>\$ 3,126</u>	<u>\$ 516</u>	<u>\$ —</u>	<u>\$ 17,805</u>
	2010					
	Security Division	Healthcare Division	Optoelectronics and Manufacturing Division	Corporate	Eliminations	Consolidated
Revenues:						
External customer revenue	\$ 251,479	\$ 206,557	\$ 137,075	\$ —	\$ —	\$ 595,111
Revenue between product segments	—	—	34,162	—	(34,162)	—
Total revenues	<u>\$ 251,479</u>	<u>\$ 206,557</u>	<u>\$ 171,237</u>	<u>\$ —</u>	<u>\$ (34,162)</u>	<u>\$ 595,111</u>
Income (loss) from operations	<u>\$ 23,351</u>	<u>\$ 13,113</u>	<u>\$ 11,958</u>	<u>\$ (10,878)</u>	<u>\$ (776)</u>	<u>\$ 36,768</u>
Segments assets	<u>\$ 221,019</u>	<u>\$ 138,739</u>	<u>\$ 85,170</u>	<u>\$ 72,731</u>	<u>\$ (4,545)</u>	<u>\$ 513,114</u>
Capital expenditures	<u>\$ 12,362</u>	<u>\$ 2,290</u>	<u>\$ 2,414</u>	<u>\$ 1,021</u>	<u>\$ —</u>	<u>\$ 18,087</u>
Depreciation and amortization	<u>\$ 6,210</u>	<u>\$ 8,365</u>	<u>\$ 3,449</u>	<u>\$ 537</u>	<u>\$ —</u>	<u>\$ 18,561</u>

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The following tables present the revenues and identifiable assets by geographical area (in thousands):

	2008				Total Consolidated
	North America	Europe	Asia	Eliminations	
Revenues:					
External customer revenue	\$ 387,655	\$ 178,281	\$ 57,152	\$ —	\$ 623,088
Revenue between product segments	27,251	—	19,816	(47,067)	—
Total revenue	<u>\$ 414,906</u>	<u>\$ 178,281</u>	<u>\$ 76,968</u>	<u>\$ (47,067)</u>	<u>\$ 623,088</u>
Long-lived assets	<u>\$ 109,478</u>	<u>\$ 29,675</u>	<u>\$ 8,788</u>		<u>\$ 147,941</u>
	2009				Total Consolidated
	North America	Europe	Asia	Eliminations	
Revenues:					
External customer revenue	\$ 397,371	\$ 135,173	\$ 57,817	\$ —	\$ 590,361
Revenue between product segments	27,530	—	18,411	(45,941)	—
Total revenue	<u>\$ 424,901</u>	<u>\$ 135,173</u>	<u>\$ 76,228</u>	<u>\$ (45,941)</u>	<u>\$ 590,361</u>
Long-lived assets	<u>\$ 108,801</u>	<u>\$ 27,520</u>	<u>\$ 11,442</u>		<u>\$ 147,763</u>
	2010				Total Consolidated
	North America	Europe	Asia	Eliminations	
Revenues:					
External customer revenue	\$ 418,808	\$ 126,450	\$ 49,853	\$ —	\$ 595,111
Revenue between product segments	23,204	—	10,958	(34,162)	—
Total revenue	<u>\$ 442,012</u>	<u>\$ 126,450</u>	<u>\$ 60,811</u>	<u>\$ (34,162)</u>	<u>\$ 595,111</u>
Long-lived assets	<u>\$ 125,485</u>	<u>\$ 23,555</u>	<u>\$ 11,289</u>		<u>\$ 160,329</u>

* * * * *

SCHEDULE II—VALUATION AND QUALIFYING ACCOUNTS

(In thousands)

<u>Description</u>	<u>Balance at Beginning of period</u>	<u>Additions</u>		<u>Deductions-Write-offs</u>	<u>Balance at end of period</u>
		<u>Charged to costs and expenses</u>	<u>Charged in other accounts</u>		
Balance for doubtful accounts:					
Year ended June 30, 2008	\$ 2,002	\$ 1,364	\$ —	\$ 1,063	\$ 2,303
Year ended June 30, 2009	\$ 2,303	\$ 5,740	\$ —	\$ 1,147	\$ 6,896
Year ended June 30, 2010	\$ 6,896	\$ 522	\$ —	\$ 1,412	\$ 6,006
Balance for warranty reserve:					
Year ended June 30, 2008	\$ 7,443	\$ 7,709	\$ —	\$ 3,555	\$ 11,597
Year ended June 30, 2009	\$ 11,597	\$ 4,472	\$ —	\$ 5,963	\$ 10,106
Year ended June 30, 2010	\$ 10,106	\$ 6,653	\$ —	\$ 5,829	\$ 10,930

INDEX TO EXHIBITS

<u>No.</u>	<u>EXHIBIT DESCRIPTION</u>
3.1	Certificate of Incorporation of OSI Systems, Inc. (1)
3.2	Certificate of Merger of OSI Systems, Inc. (1)
3.3	Bylaws of OSI Systems, Inc. (1)
4.1	Form of Common Stock Certificate (1)
10.1	Lease (A) dated June 24, 2002, between S/I Sammamish I, LLC and Spacelabs Medical, Inc. (2)
10.2	Lease (B) dated June 24, 2002, between S/I Sammamish I, LLC and Spacelabs Medical, Inc. (2)
10.3	First Amendment to Lease (A) dated October 12, 2004, between S/I Sammamish I, LLC and OSI Systems, Inc. (3)
10.4	First Amendment to Lease (B) dated October 12, 2004, between S/I Sammamish I, LLC and OSI Systems, Inc. (3)
10.5	Credit Agreement dated July 27, 2007, between Wachovia Bank, N.A. and OSI Systems, Inc. (4)
10.6	Amended and Restated 2006 Equity Participation Plan of OSI Systems, Inc. (5)
10.7	Deferred Compensation Plan of OSI Systems, Inc. (6)
10.8	First Amendment to Credit Agreement dated June 17, 2008, between Wachovia Bank, N.A. and OSI Systems, Inc. (7)
10.9	Employment Agreement dated September 22, 2008 between Alan Edrick and OSI Systems, Inc. (8)
10.10	Employment Agreement dated September 22, 2008 between Victor Sze and OSI Systems, Inc. (8)
10.11	OSI Systems, Inc. Nonqualified Defined Benefit Plan (9)
10.12	OSI Systems, Inc. 2008 Employee Stock Purchase Plan (10)
10.13	Second Amendment to Credit Agreement dated November 13, 2008, between Wachovia Bank, N.A. and OSI Systems, Inc. (11)
10.14	Employment Agreement dated September 14, 2009 between Ajay Mehra and OSI Systems, Inc. (12)
10.15	Employment Agreement dated January 6, 2010 between Deepak Chopra and OSI Systems, Inc. (13)
10.16	Third Amendment to Credit Agreement dated December 16, 2009, between Wachovia Bank, N.A. and OSI Systems, Inc. (14)
10.17*	Form of Indemnification Agreement for Directors and Executive Officers of OSI Systems, Inc.
14.1	Code of Ethics and Conduct (3)
21.1*	Subsidiaries of the Company
23.1*	Consent of Independent Registered Public Accounting Firm
31.1*	Certification Pursuant to Section 302
31.2*	Certification Pursuant to Section 302
32.1*	Certification Pursuant to Section 906
32.2*	Certification Pursuant to Section 906

* Filed herewith

(1) Previously filed with our Current Report on Form 8-K filed March 8, 2010.

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- (2) Previously filed with our Annual Report on Form 10-K filed September 13, 2004.
- (3) Previously filed with our Annual Report on Form 10-K filed September 29, 2005.
- (4) Previously filed with our Amendment No. 2 to Current Report on Form 8-K/A filed on April 13, 2010.
- (5) Previously filed with our Current Report on Form 8-K filed December 7, 2007.
- (6) Previously filed with our Current Report on Form 8-K filed June 16, 2008.
- (7) Previously filed with our Current Report on Form 8-K filed June 18, 2008.
- (8) Previously filed with our Current Report on Form 8-K filed September 23, 2008.
- (9) Previously filed with our Current Report on Form 8-K filed October 10, 2008.
- (10) Previously filed with our Proxy Statement on Schedule 14A filed October 14, 2008.
- (11) Previously filed with our Quarterly Report on Form 10-Q filed January 29, 2009.
- (12) Previously filed with our Current Report on Form 8-K filed September 16, 2009.
- (13) Previously filed with our Current Report on Form 8-K filed January 8, 2010.
- (14) Previously filed with our Quarterly Report on Form 10-Q filed January 28, 2010.

INDEMNIFICATION AGREEMENT

THIS AGREEMENT is entered into, effective as _____, _____ by and between OSI Systems, Inc., a Delaware corporation (the “Company”), and (“Indemnitee”).

A. WHEREAS, it is essential to the Company to retain and attract as directors and officers the most capable persons available;

B. WHEREAS, Indemnitee is a director and/or officer of the Company;

C. WHEREAS, both the Company and Indemnitee recognize the increased risk of litigation and other claims currently being asserted against directors and officers of corporations;

D. WHEREAS, the Certificate of Incorporation and Bylaws of the Company require the Company to indemnify and advance expenses to its directors and officers to the fullest extent permitted under Delaware law, and the Indemnitee has been serving and continues to serve as a director and/or officer of the Company in part in reliance on the Company’s Certificate of Incorporation and Bylaws; and

E. WHEREAS, in recognition of Indemnitee’s need for (i) substantial protection against personal liability based on Indemnitee’s reliance on the aforesaid Certificate of Incorporation and Bylaws, (ii) specific contractual assurance that the protection promised by the Certificate of Incorporation and Bylaws will be available to Indemnitee (regardless of, among other things, any amendment to or revocation of the Certificate of Incorporation and Bylaws or any change in the composition of the Company’s Board of Directors or acquisition transaction relating to the Company), and (iii) an inducement to provide effective services to the Company as a director and/or officer, the Company wishes to provide in this Agreement for the indemnification of and the advancing of expenses to Indemnitee to the fullest extent (whether partial or complete) permitted under Delaware law and as set forth in this Agreement, and, to the extent insurance is maintained, to provide for the continued coverage of Indemnitee under the Company’s directors’ and officers’ liability insurance policies.

NOW, THEREFORE, in consideration of the above premises and of Indemnitee continuing to serve the Company directly or, at its request, with another enterprise, and intending to be legally bound hereby, the parties agree as follows:

1. Certain Definitions.

1.1 **Board:** the Board of Directors of the Company.

1.2 **Affiliate:** any corporation or other person or entity that directly, or indirectly through one or more intermediaries, controls or is controlled by, or is under common control with, the person specified.

1.3 **Change in Control:** shall be deemed to have occurred if (i) any “person” (as such term is used in Sections 13(d) and 14(d) of the Securities Exchange Act of 1934, as amended (the “Exchange Act”)) (other than a trustee or other fiduciary holding securities under an employee benefit plan of the Company or a corporation owned directly or indirectly by the stockholders of the Company in substantially the same proportions as their ownership of stock of the Company, and other than any person holding shares of the Company on the date that the Company first registers under the Act or any transferee of such individual if such transferee is a spouse or lineal descendant of the transferee or a trust for the benefit of the individual, his spouse or lineal descendants), is or becomes the “beneficial owner” (as defined in Rule 13d-3 under the Exchange Act), directly or indirectly, of securities of the Company representing 30% or more of the total voting power represented by the Company’s then outstanding Voting Securities, or (ii) during any period of two consecutive years, individuals who at the beginning of such period constitute the Board and any new director whose election by the Board or nomination for election by the Company’s stockholders was approved by a vote of at least two-thirds (2/3) of the directors then still in office who either were directors at the beginning of the period or whose election or nomination for election was previously so approved, cease for any reason to constitute a majority of the Board, or (iii) the

stockholders of the Company approve a merger or consolidation of the Company with any other corporation, other than a merger or consolidation that would result in the Voting Securities of the Company outstanding immediately prior thereto continuing to represent (either by remaining outstanding or by being converted into Voting Securities of the surviving entity) at least 80% of the total voting power represented by the Voting Securities of the Company or such surviving entity outstanding immediately after such merger or consolidation, or (iv) the stockholders of the Company approve a plan of complete liquidation of the Company or an agreement for the sale or disposition by the Company (in one transaction or a series of transactions) of all or substantially all of the Company's assets.

1.4 **Expenses:** any expense, liability, or loss, including attorneys' fees, judgments, fines, ERISA excise taxes and penalties, amounts paid or to be paid in settlement, any interest, assessments, or other charges imposed thereon, any federal, state, local, or foreign taxes imposed as a result of the actual or deemed receipt of any payments under this Agreement, and all other costs and obligations, paid or incurred in connection with investigating, defending, being a witness in, participating in (including on appeal), or preparing for any of the foregoing in, any Proceeding relating to any Indemnifiable Event.

1.5 **Indemnifiable Event:** any event or occurrence that takes place either prior to or after the execution of this Agreement, related to the fact that Indemnitee is or was a director or officer of the Company, or while a director or officer is or was serving at the request of the Company as a director, officer, employee, trustee, agent, or fiduciary of another foreign or domestic corporation, partnership, joint venture, employee benefit plan, trust, or other enterprise, or was a director, officer, employee, or agent of a foreign or domestic corporation that was a predecessor corporation of the Company or of another enterprise at the request of such predecessor corporation, or related to anything done or not done by Indemnitee in any such capacity, whether or not the basis of the Proceeding is alleged action in an official capacity as a director, officer, employee, or agent or in any other capacity while serving as a director, officer, employee, or agent of the Company, as described above.

1.6 **Independent Counsel:** the person or body appointed in connection with Section 3.

1.7 **Proceeding:** any threatened, pending, or completed action, suit, or proceeding or any alternative dispute resolution mechanism (including an action by or in the right of the Company), or any inquiry, hearing, or investigation, whether conducted by the Company or any other party, that Indemnitee in good faith believes might lead to the institution of any such action, suit, or proceeding, whether civil, criminal, administrative, investigative, or other.

1.8 **Reviewing Party:** the person or body appointed in accordance with Section 3.

1.9 **Voting Securities:** any securities of the Company that vote generally in the election of directors.

2. Agreement to Indemnify.

2.1 **General Agreement.** In the event Indemnitee was, is, or becomes a party to or witness or other participant in, or is threatened to be made a party to or witness or other participant in, a Proceeding by reason of (or arising in part out of) an Indemnifiable Event, the Company shall indemnify Indemnitee from and against any and all Expenses to the fullest extent permitted by law, as the same exists or may hereafter be amended or interpreted (but in the case of any such amendment or interpretation, only to the extent that such amendment or interpretation permits the Company to provide broader indemnification rights than were permitted prior thereto). The parties hereto intend that this Agreement shall provide for indemnification in excess of that expressly permitted by statute, including, without limitation, any indemnification provided by the Company's Certificate of Incorporation, its Bylaws, vote of its shareholders or disinterested directors, or applicable law.

2.2 **Initiation of Proceeding.** Notwithstanding anything in this Agreement to the contrary, Indemnitee shall not be entitled to indemnification pursuant to this Agreement in connection with any Proceeding initiated by Indemnitee against the Company or any director or officer of the Company unless (i) the Company has joined in or the Board has consented to the initiation of such Proceeding; (ii) the Proceeding is one to enforce indemnification rights under Section 5; or (iii) the Proceeding is instituted after a Change in Control and Independent Counsel has approved its initiation.

2.3 **Expense Advances.** If so requested by Indemnitee, whether in a single instance or from time to time, the Company shall advance (within ten business days of such request) any and all Expenses to Indemnitee (an "Expense Advance"); provided that, (i) such an Expense Advance shall be made only upon delivery to the Company of an undertaking by or on behalf of the Indemnitee to repay the amount thereof if it is ultimately determined that Indemnitee is not entitled to be indemnified by the Company, and (ii) if and to the extent that the Reviewing Party determines that Indemnitee would not be permitted to be so indemnified under applicable law, the Company shall be entitled to be reimbursed by Indemnitee (who hereby agrees to reimburse the Company) for all such amounts theretofore paid. If Indemnitee has commenced or commences legal proceedings in a court of competent jurisdiction to secure a determination that Indemnitee should be indemnified under applicable law, as provided in Section 4, any determination made by the Reviewing Party that Indemnitee would not be permitted to be indemnified under applicable law shall not be binding, and Indemnitee shall not be required to reimburse the Company for any Expense Advance until a final judicial determination is made with respect thereto (as to which all rights of appeal therefrom have been exhausted or have lapsed); while any such final judicial determination is pending, the Company shall be obligated to provide Expense Advances. Indemnitee's obligation to reimburse the Company for Expense Advances shall be unsecured and no interest shall be charged thereon.

2.4 **Mandatory Indemnification.** Notwithstanding any other provision of this Agreement, to the extent that Indemnitee has been successful on the merits or otherwise in defense of any Proceeding relating in whole or in part to an Indemnifiable Event or in defense of any claim, issue or matter therein, Indemnitee shall be indemnified against all Expenses actually and reasonably incurred by Indemnitee in connection therewith.

2.5 **Partial Indemnification.** If Indemnitee is entitled under any provision of this Agreement to indemnification by the Company for some or a portion of Expenses, but not, however, for the total amount thereof, the Company shall nevertheless indemnify Indemnitee for the portion thereof to which Indemnitee is entitled.

2.6 **Prohibited Indemnification.** No indemnification pursuant to this Agreement shall be paid by the Company on account of any Proceeding in which judgment is rendered against Indemnitee for an accounting of profits made from the purchase or sale by Indemnitee of securities of the Company pursuant to the provisions of Section 16(b) of the Securities Exchange Act of 1934, as amended, or similar provisions of any federal, state, or local laws.

3. **Reviewing Party.** Prior to any Change in Control, the Reviewing Party shall be (a) the Board, acting by a majority vote of the directors who are not parties to the Proceeding, even though less than a quorum of the Board, (b) a committee of directors designated by a majority vote of the directors, even though less than a quorum of the Board, (c) if there are no such directors or, if such directors so direct, Independent Counsel, or (d) if so directed by the Board, the stockholders of the Company; after a Change in Control, the Independent Counsel referred to below shall become the Reviewing Party. With respect to all matters arising after a Change in Control concerning the rights of Indemnitee to indemnity payments and Expense Advances under this Agreement or any other agreement or under applicable law or the Company's Certificate of Incorporation or Bylaws now or hereafter in effect relating to indemnification for Indemnifiable Events, the Company shall seek legal advice only from Independent Counsel selected by Indemnitee and approved by the Company (which approval shall not be unreasonably withheld), and who has not otherwise performed services for the Company or the Indemnitee (other than in connection with indemnification matters) within the last five years. The Independent Counsel shall not include any person who, under the applicable standards of professional conduct then prevailing, would have a conflict of interest in representing either the Company or Indemnitee in an action to determine Indemnitee's rights under this Agreement. Such counsel, among other things, shall render its written opinion to the Company and Indemnitee as to whether and to what extent the Indemnitee should be permitted to be indemnified under applicable law. The Company agrees to pay the reasonable fees of the Independent Counsel and to indemnify fully such counsel against any and all expenses (including attorneys' fees), claims, liabilities, loss, and damages arising out of or relating to this Agreement or the engagement of Independent Counsel pursuant hereto.

4. Indemnification Process and Appeal.

4.1 **Indemnification Payment.** Indemnitee shall be entitled to indemnification of Expenses, and shall receive payment thereof, from the Company in accordance with this Agreement as soon as practicable after Indemnitee has made written demand on the Company for indemnification, unless the Reviewing Party has given a written opinion to the Company that Indemnitee is not entitled to indemnification under applicable law.

4.2 **Suit to Enforce Rights.** Regardless of any action by the Reviewing Party, if Indemnitee has not received full indemnification within thirty days after making a demand in accordance with Section 4.1, Indemnitee shall have the right to enforce its indemnification rights under this Agreement by commencing litigation in the Delaware Chancery Court or any court located in the County of Los Angeles having subject matter jurisdiction thereof seeking an initial determination by the court or challenging any determination by the Reviewing Party or any aspect thereof. The Company hereby consents to service of process and to appear in any such proceeding. Any determination by the Reviewing Party not challenged by the Indemnitee shall be binding on the Company and Indemnitee. The remedy provided for in this Section 4 shall be in addition to any other remedies available to Indemnitee at law or in equity. Any suit initiated by the Company with respect to this Agreement shall be filed in the Delaware Chancery Court or any court located in the County of Los Angeles.

4.3 **Defense to Indemnification, Burden of Proof, and Presumptions.** It shall be a defense to any action brought by Indemnitee against the Company to enforce this Agreement (other than an action brought to enforce a claim for Expenses incurred in defending a Proceeding in advance of its final disposition where the required undertaking has been tendered to the Company) that it is not permissible under applicable law for the Company to indemnify Indemnitee for the amount claimed. In connection with any such action or any determination by the Reviewing Party or otherwise as to whether Indemnitee is entitled to be indemnified hereunder, the burden of proving such a defense or determination shall be on the Company. Neither the failure of the Reviewing Party or the Company (including its Board, independent legal counsel, or its stockholders) to have made a determination prior to the commencement of such action by Indemnitee that indemnification of the claimant is proper under the circumstances because Indemnitee has met the standard of conduct set forth in applicable law, nor an actual determination by the Reviewing Party or Company (including its Board, independent legal counsel, or its stockholders) that the Indemnitee had not met such applicable standard of conduct, shall be a defense to the action or create a presumption that the Indemnitee has not met the applicable standard of conduct. For purposes of this Agreement, the termination of any claim, action, suit, or proceeding, by judgment, order, settlement (whether with or without court approval), conviction, or upon a plea of nolo contendere, or its equivalent, shall not create a presumption that Indemnitee did not meet any particular standard of conduct or have any particular belief or that a court has determined that indemnification is not permitted by applicable law.

5. **Indemnification for Expenses Incurred in Enforcing Rights.** The Company shall indemnify Indemnitee against any and all Expenses that are incurred by Indemnitee in connection with any action brought by Indemnitee for (i) indemnification or advance payment of Expenses by the Company under this Agreement or any other agreement or under applicable law or the Company's Certificate of Incorporation or Bylaws now or hereafter in effect relating to indemnification for Indemnifiable Events, and/or (ii) recovery under directors' and officers' liability insurance policies maintained by the Company, but only in the event that Indemnitee ultimately is determined to be entitled to such indemnification or insurance recovery, as the case may be. In addition, the Company shall, if so requested by Indemnitee, advance the foregoing Expenses to Indemnitee, subject to and in accordance with Section 2.3.

6. Notification and Defense of Proceeding.

6.1 **Notice.** Promptly after receipt by Indemnitee of notice of the commencement of any Proceeding, Indemnitee shall, if a claim in respect thereof is to be made against the Company under this Agreement, notify the Company of the commencement thereof; but the omission so to notify the Company will not relieve the Company from any liability that it may have to Indemnitee, except as provided in Section 6.3.

6.2 **Defense.** With respect to any Proceeding as to which Indemnitee notifies the Company of the commencement thereof, the Company will be entitled to participate in the Proceeding at its own expense and

except as otherwise provided below, to the extent the Company so wishes, it may assume the defense thereof with counsel reasonably satisfactory to Indemnitee. After notice from the Company to Indemnitee of its election to assume the defense of any Proceeding (including costs thereof), the Company shall not be liable to Indemnitee under this Agreement or otherwise for any other Expenses subsequently incurred by Indemnitee in connection with the defense of such Proceeding except for reasonable costs of investigation or as otherwise provided below. Indemnitee shall have the right to employ independent legal counsel in such Proceeding, but all Expenses related thereto incurred after notice from the Company of its assumption of the defense shall be at Indemnitee's expense unless any one or more of the following occur: (i) the employment of legal counsel by Indemnitee has been authorized by the Company, (ii) Indemnitee has reasonably determined that there may be a conflict of interest between Indemnitee and the Company in the defense of the Proceeding, (iii) after a Change in Control, the employment of counsel by Indemnitee has been approved by the Independent Counsel, or (iv) the Company shall not in fact have employed counsel to assume the defense of such Proceeding, in each of which cases ((i) through (iv)) all Expenses of the Proceeding shall be borne by the Company. The Company shall not be entitled to assume the defense of any Proceeding brought by or on behalf of the Company or as to which Indemnitee shall have made the determination provided for in (ii), (iii) and (iv) above.

6.3 **Settlement of Claims.** The Company shall not be liable to indemnify Indemnitee under this Agreement or otherwise for any amounts paid in settlement of any Proceeding effected without the Company's written consent, such consent not to be unreasonably withheld; provided, however, that if a Change in Control has occurred, the Company shall be liable for indemnification of Indemnitee for amounts paid in settlement if the Independent Counsel has approved the settlement. The Company shall not settle any Proceeding in any manner that would impose any penalty or limitation on Indemnitee without Indemnitee's written consent. The Company shall not be liable to indemnify the Indemnitee under this Agreement with regard to any judicial award if the Company was not given a reasonable and timely opportunity, at its expense, to participate in the defense of such action; the Company's liability hereunder shall not be excused if participation in the Proceeding by the Company was barred by this Agreement.

7. **Non-Exclusivity.** The rights of Indemnitee hereunder shall be in addition to any other rights Indemnitee may have under the Company's Certificate of Incorporation, Bylaws, applicable law, or otherwise; provided, however, that this Agreement shall supersede any prior indemnification agreement between the Company and the Indemnitee. To the extent that a change in applicable law (whether by statute or judicial decision) permits greater indemnification than would be afforded currently under the Company's Certificate of Incorporation, Bylaws, applicable law, or this Agreement, it is the intent of the parties that Indemnitee enjoy by this Agreement the greater benefits so afforded by such change.

8. **Liability Insurance.** To the extent the Company maintains an insurance policy or policies providing general and/or directors' and officers' liability insurance, Indemnitee shall be covered by such policy or policies, in accordance with its or their terms, to the maximum extent of the coverage available for any Company director or officer.

9. **Period of Limitations.** No legal action shall be brought and no cause of action shall be asserted by or on behalf of the Company or any Affiliate of the Company against Indemnitee, Indemnitee's spouse, heirs, executors, or personal or legal representatives after the expiration of two years from the date of accrual of such cause of action, or such longer period as may be required by state law under the circumstances. Any claim or cause of action of the Company or its Affiliate shall be extinguished and deemed released unless asserted by the timely filing and notice of a legal action within such period; provided, however, that if any shorter period of limitations is otherwise applicable to any such cause of action, the shorter period shall govern.

10. **Amendment of this Agreement.** No supplement, modification, or amendment of this Agreement shall be binding unless executed in writing by both of the parties hereto. No waiver of any of the provisions of this Agreement shall be binding unless in the form of a writing signed by the party against whom enforcement of the waiver is sought, and no such waiver shall operate as a waiver of any other provisions hereof (whether or not similar), nor shall such waiver constitute a continuing waiver. Except as specifically provided herein, no failure to exercise or any delay in exercising any right or remedy hereunder shall constitute a waiver thereof.

11. **Subrogation.** In the event of payment under this Agreement, the Company shall be subrogated to the extent of such payment to all of the rights of recovery of Indemnitee, who shall execute all papers required and shall do everything that may be necessary to secure such rights, including the execution of such documents necessary to enable the Company effectively to bring suit to enforce such rights.

12. **No Duplication of Payments.** The Company shall not be liable under this Agreement to make any payment in connection with any claim made against Indemnitee to the extent Indemnitee has otherwise received payment (under any insurance policy, Bylaw, or otherwise) of the amounts otherwise indemnifiable hereunder.

13. **Binding Effect.** This Agreement shall be binding upon and inure to the benefit of and be enforceable by the parties hereto and their respective successors (including any direct or indirect successor by purchase, merger, consolidation, or otherwise to all or substantially all of the business and/or assets of the Company), assigns, spouses, heirs, and personal and legal representatives. The Company shall require and cause any successor (whether direct or indirect by purchase, merger, consolidation, or otherwise) to all, substantially all, or a substantial part, of the business and/or assets of the Company, by written agreement in form and substance satisfactory to Indemnitee, expressly to assume and agree to perform this Agreement in the same manner and to the same extent that the Company would be required to perform if no such succession had taken place. The indemnification provided under this Agreement shall continue as to Indemnitee for any action taken or not taken while serving in an indemnified capacity pertaining to an Indemnifiable Event even though he may have ceased to serve in such capacity at the time of any Proceeding.

14. **Severability.** If any provision (or portion thereof) of this Agreement shall be held by a court of competent jurisdiction to be invalid, void, or otherwise unenforceable, the remaining provisions shall remain enforceable to the fullest extent permitted by law. Furthermore, to the fullest extent possible, the provisions of this Agreement (including, without limitation, each portion of this Agreement containing any provision held to be invalid, void, or otherwise unenforceable, that is not itself invalid, void, or unenforceable) shall be construed so as to give effect to the intent manifested by the provision held invalid, void, or unenforceable.

15. **Governing Law.** This Agreement shall be governed by and construed and enforced in accordance with the laws of the State of Delaware applicable to contracts made and to be performed in such State without giving effect to its principles of conflicts of laws.

16. **Notices.** All notices, demands, and other communications required or permitted hereunder shall be made in writing and shall be deemed to have been duly given if delivered by hand, against receipt, or mailed, postage prepaid, certified or registered mail, return receipt requested, and addressed to the Company at:

OSI Systems, Inc.
ATTN: Chief Executive Officer
12525 Chadron Avenue
Hawthorne, CA 90250

With a copy to:

OSI Systems, Inc.
ATTN: General Counsel
12525 Chadron Avenue
Hawthorne, CA 90250
and to Indemnitee at:

Notice of change of address shall be effective only when given in accordance with this Section. All notices complying with this Section shall be deemed to have been received on the date of hand delivery or on the third business day after mailing.

17. **Counterparts.** This Agreement may be executed in one or more counterparts, each of which shall be deemed an original, but all of which together shall constitute one and the same instrument.

IN WITNESS WHEREOF, the parties hereto have duly executed and delivered this Agreement as of the day specified above.

OSI SYSTEMS, INC.

By: _____

Its: _____

INDEMNITEE

SUBSIDIARIES OF OSI SYSTEMS, INC.

Blease Medical Holdings Ltd	London, United Kingdom
CXR Limited	Surrey, United Kingdom
Del Mar Reynolds Medical Ltd	London, United Kingdom
Dolphin Medical Pte Ltd.	Singapore
Dolphin Medical, Inc.	Hawthorne, California
ECIL Rapiscan Ltd.	Andhra Pradesh, India
Ferson Technologies, Inc.	Ocean Springs, Mississippi
Metorex Security Products, Inc.	Torrance, California
Opto Sensors Hong Kong Limited	Hong Kong
OSI Defense Systems, L.L.C.	Orlando, Florida
OSI Electronics Pte Ltd	Singapore
OSI Electronics, Inc.	Camarillo, California
OSI Optoelectronics AS	Horten, Norway
OSI Optoelectronics Limited	Nicosia, Cyprus
OSI Optoelectronics, Inc.	Hawthorne, California
OSI Optoelectronics Sdn. Bhd.	Johor Bahru, Malaysia
OSI Systems Private Limited	Andhra Pradesh, India
Osteometer MediTech, Inc.	Hawthorne, California
PT OSI Electronics	Batam, Indonesia
PT OSI Systems	Batam, Indonesia
Rapiscan Systems Electrical Trading LLC	Abu Dhabi
Rapiscan do Brazil Ltda	Rio de Janeiro, Brazil
Rapiscan Laboratories, Inc.	Sunnyvale, California
Rapiscan Systems (Cyprus) Limited	Nicosia, Cyprus
Rapiscan Systems Australia Pty Ltd	Victoria, Australia
Rapiscan Systems Hong Kong Limited	Hong Kong
Rapiscan Systems Limited	Salfords, United Kingdom
Rapiscan Systems Oy	Espoo, Finland
Rapiscan Systems Pte. Ltd.	Singapore
Rapiscan Systems Sdn. Bhd.	Johor Bahru, Malaysia
Rapiscan Systems, Inc.	Hawthorne, California
Rapiscan Systems, S.A. de C.V.	Colonia Juarez, Mexico
S2 Global, Inc.	Delaware
S2 Services Puerto Rico, LLC	Puerto Rico
S2 Services, Ltd.	Cayman Islands
SL Healthcare Limited	Nicosia, Cyprus

Spacelabs Healthcare (Canada), Inc.

Spacelabs Healthcare GmbH

Spacelabs Healthcare, Inc.

Spacelabs Healthcare, LLC

Spacelabs Healthcare Ltd.

Spacelabs Healthcare Medical Equipment (Suzhou) Co., Ltd

Spacelabs Healthcare Pte. Ltd.

Spacelabs Healthcare SAS

Spacelabs Healthcare s.r.l.

Spacelabs Healthcare Trading (Shanghai) Co., Ltd

Spacelabs Medical, Inc.

Ontario, Canada

Feucht, Germany

Delaware

Issaquah, Washington

United Kingdom

Suzhou, China

Singapore

Creteil, France

Verona, Italy

Shanghai, China

Delaware

CONSENT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Board of Directors and Shareholders
OSI Systems, Inc.

We consent to the inclusion in this Annual Report on Form 10-K of OSI Systems, Inc. for the year ended June 30, 2010 and to the incorporation by reference in Registration Statements on Forms S-8 (No. 333-45049, 333-106176, 333-122674, 333-69433, 333-132142, 333-148936 and 333-157032) and in Registration Statements on Forms S-3 (No. 333-148937, 333-119704, 333-75228, 333-73618, 333-100791 and 333-101716) of OSI Systems, Inc. of our report dated August 26, 2010 appearing in Item 8 in this Annual Report on Form 10-K, on the consolidated financial statements and financial statement schedule, which appears in Schedule II of this Form 10-K, and of our report dated August 26, 2010 with respect to the effectiveness of internal control over financial reporting, which report is included in Item 9A in this Annual Report on Form 10-K.

/s/ MOSS ADAMS LLP

Los Angeles, California
August 26, 2010

CERTIFICATION

I, Deepak Chopra, certify that:

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

Date: August 26, 2010

/s/ DEEPAK CHOPRA

Deepak Chopra
Chief Executive Officer

CERTIFICATION

I, Alan Edrick, certify that:

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

Date: August 26, 2010

/s/ ALAN EDRICK

Alan Edrick
Chief Financial Officer

**CERTIFICATION PURSUANT TO 18 U.S.C. SECTION 1350
AS ADOPTED PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002**

In connection with the Annual Report of OSI Systems, Inc. (the Company) on Form 10-K for the year ended June 30, 2010 as filed with the Securities and Exchange Commission on the date hereof (Report), I, Deepak Chopra, Chief Executive Officer of the Company, certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

- (a) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, as amended; and
- (b) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company at the dates and for the periods presented in this Report.

Date: August 26, 2010

/s/ DEEPAK CHOPRA

Deepak Chopra
Chief Executive Officer

**CERTIFICATION PURSUANT TO 18 U.S.C. SECTION 1350
AS ADOPTED PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002**

In connection with the Annual Report of OSI Systems, Inc. (the Company) on Form 10-K for the year ended June 30, 2010 as filed with the Securities and Exchange Commission on the date hereof (Report), I, Alan Edrick, Chief Financial Officer of the Company, certify, pursuant to 18, U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

(a) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, as amended; and

(b) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company at the dates and for the periods presented in this Report.

Date: August 26, 2010

/s/ ALAN EDRICK

Alan Edrick
Chief Financial Officer