
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION**
Washington, DC 20549

FORM 10-K

Annual report pursuant to section 13 or 15(d) of the Securities Exchange Act of 1934

For the fiscal year ended December 31, 2013

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 No. 001-33624

Amedica Corporation

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction of
incorporation or organization)

84-1375299
(IRS Employer
Identification No.)

1885 West 2100 South, Salt Lake City, UT 84119
(Address of principal executive offices and Zip Code)

(801) 839-3500
(Registrant's telephone number, including area code)

<u>Title of each class</u>	<u>Name of each exchange on which registered</u>
Common Stock, \$0.01 par value	The NASDAQ Capital Market

Securities registered under Section 12(b) of the Act: Common Stock, Par Value \$0.01

Securities registered under Section 12(g) of the Act:

None

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

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

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

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Website, if any, every interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). Yes No

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

Large Accelerated Filer

Accelerated Filer

Non-Accelerated Filer [Do not check if a smaller reporting company]

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 of the registrant (without admitting that any person whose shares are not included in such calculation is an affiliate) computed by reference to the initial public offering price, on February 12, 2014 was \$58,847,024. The registrant has provided this information as of February 12, 2014 because its common stock was not publicly traded as of the last business day of its most recently completed second fiscal quarter.

The number of shares outstanding of the registrant's common stock, \$0.01 par value per share, as of March 14, 2014 was 12,310,354.

DOCUMENTS INCORPORATED BY REFERENCE: NONE

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CAUTIONARY NOTE CONCERNING FORWARD-LOOKING STATEMENTS

This Annual Report on Form 10-K 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 (the “Securities Act”), and Section 21E of the Securities Exchange Act of 1934, as amended (the “Exchange Act”). All statements other than statements of historical fact are forward-looking statements. We have tried to identify forward-looking statements by using words such as “believe,” “may,” “might,” “could,” “will,” “aim,” “estimate,” “continue,” “anticipate,” “intend,” “expect,” “plan” and similar words. These forward-looking statements are based on our current assumptions, expectations and estimates of future events and trends. Forward-looking statements are only predictions and are subject to many risks, uncertainties and other factors that may affect our businesses and operations and could cause actual results to differ materially from those predicted. These risks and uncertainties include, but are not limited to, factors affecting our quarterly results, our ability to manage our growth, our ability to sustain our profitability, demand for our products, our ability to compete successfully (including without limitation our ability to convince surgeons to use our products and our ability to attract and retain sales and other personnel), our ability to rapidly develop and introduce new products, our ability to develop and execute on successful business strategies, our ability to comply with changes and applicable laws and regulations that are applicable to our businesses, our ability to safeguard our intellectual property, our success in defending legal proceedings brought against us, trends in the medical device industry, and general economic conditions, and other risks set forth throughout this Annual Report, including under “**Item 1, Business,**” “**Item 1A, Risk Factors,**” and “**Item 7, Management’s Discussion and Analysis of Financial Condition and Results of Operations,**” and those discussed in other documents we file with the Securities and Exchange Commission (the “SEC”). Moreover, we operate in an evolving environment. New risk factors and uncertainties emerge from time to time and it is not possible for us to predict all risk factors and uncertainties, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statements.

Given these risks and uncertainties, readers are cautioned not to place undue reliance on any forward-looking statements. Forward-looking statements contained in this Annual Report speak only as of the date of this Annual Report. We undertake no obligation to update any forward-looking statements as a result of new information, events or circumstances or other factors arising or coming to our attention after the date hereof.

WHERE YOU CAN FIND MORE INFORMATION

We are subject to the informational requirements of the Exchange Act. Accordingly, we file periodic reports and other information with the SEC. We will make our annual report on Form 10-K, quarterly reports on Form 10-Q, current reports on Form 8-K and amendments to those reports available through our Internet site, http://www.amedica.com/investors/sec_filings as soon as reasonably practicable after electronically filing such materials with the SEC. They may also be obtained free of charge by writing to Amedica Corporation, Attn: Investor Relations, 1885 West 2100 South, Salt Lake City, UT 84119. In addition, copies of these reports may be obtained through the SEC’s website at www.sec.gov or by visiting the SEC’s Public Reference Room at 100 F Street, NE, Washington, DC 20549 or by calling the SEC at 800-SEC-0330. Our common stock trades on The NASDAQ Capital Market under the symbol “AMDA.”

Unless otherwise indicated, all information contained in this Annual Report reflects a 1-for-25.7746 reverse split of our common stock which was effected on February 11, 2014.

PART I

ITEM 1. BUSINESS

Overview

We are a commercial biomaterial company focused on using our silicon nitride technology platform to develop, manufacture and sell a broad range of medical devices. We currently market spinal fusion products and are developing products for use in total hip and knee joint replacements. We believe our silicon nitride, an advanced ceramic, technology platform enables us to offer new and transformative products in the orthopedic and other medical device markets. We believe we are the first and only company to use silicon nitride in medical applications and over 14,000 of our silicon nitride spine products have been implanted in patients.

Biomaterials are synthetic or natural materials available in a variety of forms that are used in virtually every medical specialty. We believe our silicon nitride biomaterial has superior characteristics compared to commonly used biomaterials in the markets we are targeting, including polyetheretherketone, or PEEK, which is the most common biomaterial used for interbody spinal fusion products. Specifically, we believe our silicon nitride has the following key attributes: promotion of bone growth; hardness, strength and resistance to fracture; resistance to wear; non-corrosive properties; anti-infective properties; and superior diagnostic imaging compatibility.

We produce our silicon nitride advanced ceramic in four forms: (1) a fully dense, load-bearing solid, referred to as *MC²*; (2) a porous bone-like cancellous structured form, referred to as *C^{SC}*; (3) a composite incorporating both our solid *MC²* material and our porous *C^{SC}* material intended to promote an ideal environment for bone growth; and (4) a coating for application onto other biomaterials. This capability provides us with the ability to utilize our silicon nitride in distinct ways depending on its intended application, which, together with our silicon nitride's key characteristics, distinguishes us from manufacturers of other biomaterials and our products from products using other biomaterials.

According to iData Research, Inc., or iData, in 2012, the markets for spinal implants in the United States and in combined major European markets were \$5.2 billion and \$1.0 billion, respectively. Interbody spinal fusions accounted for over \$1.2 billion and \$172.2 million of these markets, respectively. Additionally, Orthopedic Network News reported that the U.S. markets for the components of total hip and knee replacement product candidates that we are initially developing were \$455.0 million and \$1.5 billion, respectively.

We currently market our *Valeo MC²* silicon nitride interbody spinal fusion devices in the United States and Europe for use in the cervical and thoracolumbar areas of the spine. We believe our *Valeo* devices have a number of advantages over existing products due to silicon nitride's key characteristics, resulting in faster and more effective fusion and reduced risk of infection. Our first generation *Valeo* silicon nitride device received 510(k) regulatory clearance and a CE Mark in 2008. Based on surgeon feedback, we developed a second generation of *Valeo* AL, PL and OL products with design enhancements that improve surgeon control during implantation and stability post procedure. In 2013, we initiated a targeted launch of our second generation AL, PL and OL *Valeo* interbody fusion devices and expect to complete the full launch in the second quarter of 2014. We are also completing the development of our second generation Cervical and TL *Valeo* interbody spinal fusion devices and expect these to be launched in the second half of 2014. We also market our *Valeo* composite interbody spinal fusion device made from both our solid *MC²* and porous *C^{SC}* silicon nitride in the Netherlands, Spain and Germany. This device may reduce or eliminate the need for allograft bone, which is taken from human cadavers, and other biomaterials to act as a scaffold to support bone growth as part of the surgical procedure. We are currently conducting a prospective clinical trial in Europe, named CASCADE, comparing our *Valeo* composite silicon nitride interbody devices to PEEK interbody devices to obtain additional data to support 510(k) clearance of this product in the United States. The trial is 100% enrolled. We expect results to be available in the second half of 2014. If this trial is successful, we plan to file a 510(k) submission with the U.S. Food and Drug Administration, or FDA, by mid-2015. In addition, in the first half of 2013, we initiated a Design and Build Program focused on collaborating with influential surgeons to develop customized silicon nitride spinal fusion products and instruments and the first products designed under this program were sold in the third quarter of 2013. As of December 31, 2013, the rate of adverse events reported to the FDA for our implanted *Valeo* interbody spinal fusion devices is 0.1%.

In addition to our silicon nitride-based spinal fusion products, we market a complementary line of non-silicon nitride spinal fusion products which allows us to provide surgeons and hospitals with a broader range of products. These products include three lines of spinal fusion devices and five types of orthobiologics, which are used by surgeons to help promote bone growth and fusion in spinal fusion procedures. Although our non-silicon nitride products have accounted for approximately 66% and 74% or more of our product revenues for the years ended December 31, 2013 and 2012, respectively, we believe the continued promotion and potential for adoption of our silicon nitride products and product candidates, if approved, provides us the greatest opportunity to grow our business in new and existing markets and achieve our goal to become a leading biomaterial company.

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We are also incorporating our silicon nitride technology into components for use in total hip and knee replacement product candidates that we are, or plan on, developing in collaboration with a strategic partner. If approved by the FDA, we believe that our silicon nitride total hip and knee product candidates will provide competitive advantages over current products made with traditional biomaterials. We also believe our silicon nitride technology platform can be used for developing products in other markets and have developed prototypes for use in the dental, sports medicine and trauma markets. In addition, as a result of some of the key characteristics of our silicon nitride, including the promotion of bone growth, resistance to wear, non-corrosiveness and anti-infective properties, we believe our silicon nitride coating may be used to enhance our metal products as well as commercially available metal spinal fusion, joint replacement and other medical products.

We have recently put in place a senior management team with over 150 years of collective experience in the healthcare industry. Members of our management team have experience in product development, launching new products into the orthopedics market and selling to hospitals through direct sales organizations, distributors, manufacturers and other companies in the orthopedic space. We operate a 30,000 square foot manufacturing facility located at our corporate headquarters in Salt Lake City, Utah, and we are the only vertically integrated silicon nitride orthopedic medical device manufacturer in the world. We market and sell our products to surgeons and hospitals in the United States and select markets in Europe and South America through our established network of more than 50 independent sales distributors who are managed by our in-house sales and marketing management team.

Biomaterials

Biomaterials are synthetic or natural biocompatible materials that are used in virtually every medical specialty to improve or preserve body functionality. Various types of biomaterials are used as essential components in medical devices, drug delivery systems, replacement and tissue repair technologies, prostheses and diagnostic technologies.

There are four general categories of biomaterials:

- *Metals.* Metals commonly used as biomaterials include titanium, stainless steel, cobalt, chrome, gold, silver and platinum, and alloys of these metals. Examples of medical uses of metals include the repair or stabilization of fractured bones, stents, surgical instruments, bone and joint replacements, spinal fusion devices, dental implants and restorations and heart valves. According to MarketsandMarkets, a global market research firm, metals represented approximately 31% of the worldwide sales of all biomaterials in 2012.
- *Polymers.* Polymers are synthetic compounds consisting of similar molecules linked together that can be created to have specific properties. Polymers commonly used as biomaterials include nylon, silicon rubber, polyester, polyethylene, cross-linked polyethylene (a stronger version), polymethylmethacrylate, polyvinyl chloride and polyetheretherketone, which is commonly referred to as PEEK. Examples of medical uses of polymers include soft-tissue replacement, sutures, drug delivery systems, joint replacements, spinal fusion devices and dental restorations. Polymers represented approximately 29% of the worldwide sales of all biomaterials in 2012.
- *Ceramics.* Ceramics are hard, non-metallic, non-corrosive, heat-resistant materials made by shaping and then applying high temperatures. Traditional ceramics commonly used as biomaterials include carbon, oxides of aluminum, zirconium and titanium, calcium phosphate and zirconia-toughened alumina. Examples of medical uses of ceramics include repair, augmentation or stabilization of fractured bones, bone and joint replacements, spinal fusion devices, dental implants and restorations, heart valves and surgical instruments. Ceramics represented approximately 26% of the worldwide sales of all biomaterials in 2012.
- *Natural biomaterials.* Natural biomaterials are derived from human donors, animal or plant sources and include human bone, collagen, gelatin, cellulose, chitin, alginate and hyaluronic acid. Examples of medical uses of natural biomaterials include the addition or substitution of hard and soft tissue, cornea protectors, vascular grafts, repair and replacement of tendons and ligaments, bone and joint replacements, spinal fusion devices, dental restorations and heart valves. Natural biomaterials represented approximately 14% of the worldwide sales of all biomaterials in 2012.

According to MarketsandMarkets, orthopedics accounted for approximately \$15.0 billion, or 34%, of the \$44.0 billion total biomaterials market in 2012. Within orthopedics, biomaterials are extensively used in spinal fusion procedures, hip and knee replacements and the repair or stabilization of fractured bones.

Market Opportunity

Overview

We believe our silicon nitride technology platform provides us with numerous competitive advantages in the orthopedic biomaterials market. We market interbody spinal fusion devices and related products and are developing products for use as components in total hip and knee joint replacements. We believe we can also utilize our silicon nitride technology platform to develop future products in additional markets, such as the dental, sports medicine and trauma markets.

According to iData, in 2012, the markets for spinal implants in the United States and in combined major European markets were \$5.3 billion and \$1.0 billion, respectively. Interbody spinal fusion products accounted for over \$1.2 billion and \$172.2 million of these markets, respectively. In 2012, there were approximately 300,000 interbody spinal fusion procedures conducted in the United States, of which the significant majority utilized interbody devices comprised of PEEK and bone, with occasional use of metals and other materials including ceramics. The market for interbody spinal fusion devices has shifted over time as new biomaterials with superior characteristics have been incorporated into these devices and have launched into the market. For example, in the 1990s, metals quickly penetrated the interbody spinal fusion market because of the limitations of devices available at that time made from allograft bone and, more recently, products made of PEEK rapidly penetrated the market because of the limitations of devices available at that time made from metal or allograft bone. Similarly, we believe our silicon nitride interbody spinal fusion products address the key limitations of other biomaterials currently used in interbody spinal fusion devices and demonstrate superior characteristics needed to improve clinical outcomes.

Additionally, Orthopedic Network News reported that the U.S. markets for total hip and knee replacements in 2012 were \$2.7 billion and \$4.0 billion, respectively. According to Orthopedic Network News, in 2012, there were more than 470,000 total hip replacement procedures and 734,000 total knee replacement procedures conducted in the United States. Orthopedic Network News also reported that in 2012, the U.S. markets for the components of total hip and knee replacement product candidates that we are initially developing were \$455.0 million and \$1.5 billion, respectively. The combinations of biomaterials most commonly used in joint replacement implants are metal-on-cross-linked polyethylene and traditional ceramic-on-cross-linked polyethylene.

We believe that the main drivers for the growth of the orthopedic biomaterials market, and, in particular, the spinal fusion and joint replacement markets, are the following:

- *Favorable and Changing Demographics.* With the growing number of elderly people, age-related ailments are expected to rise sharply, which we believe will increase the demand and need for biomaterials and devices with improved performance capabilities. Also, middle-aged and older patients increasingly expect to enjoy active lifestyles, and consequently demand effective treatments for painful spine and joint conditions, including better performing and longer-lasting interbody spinal fusion devices and joint replacements.
- *Introduction of New Technologies.* Better performing and longer-lasting biomaterials, improved diagnostics, and advances in surgical procedures allow for surgical intervention earlier in the continuum of care and better outcomes for patients. We believe surgical options using better performing and longer-lasting biomaterials will gain acceptance among surgeons and younger patients and drive accelerated growth and increase the size of the spinal fusion and joint replacement markets.
- *Market Expansion into New Geographic Areas.* MarketsandMarkets anticipates that demand for biomaterials and the associated medical devices will increase as the applications in which biomaterials are used are introduced to and become more widely accepted in underserved countries, such as China.

The Interbody Spinal Fusion Market

The human spinal canal is made up of 33 interlocking bones, referred to as vertebrae, separated by 23 intervertebral discs comprised of a hard outer ring made of collagen with a soft inner core, that act as shock absorbers between vertebrae. Disorders of the spine can result from degenerative conditions, deformities and trauma or tumor-related damage. Spinal fusion is the standard of care used to treat most spinal disorders and typically involves the placement of an interbody device between vertebrae to reestablish spacing between vertebrae and alignment of the spine. Generally, the interbody device is stabilized by screws and, in some procedures, plates or rods. To enhance bone attachment, surgeons often pack the interbody device with a biomaterial that induces bone growth. Following successful treatment, new bone tissue grows in and around the interbody device over time, which helps fuse the vertebrae and create long-term stability of the interbody device, leading to the alleviation of pain and increase in mobility.

We selected this market as the first application for our silicon nitride technology because of its size, the limitations of currently available products and the key characteristics silicon nitride possesses which are critical for a superior interbody spinal fusion device.

- *Promotion of Bone Growth.* The biomaterial should be both osteoconductive and create an osteoinductive environment to promote bone growth in and around the interbody device to further support fusion and stability. Osteoconduction occurs when material serves as a scaffold to support the growth of new bone in and around the material. Osteoinduction involves the stimulation of osteoprogenitor cells to develop, or differentiate, into osteoblasts, which are cells that are needed for bone growth.

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- *Strength and Resistance to Fracture.* The biomaterial should be strong and resistant to fracture during implantation of the device and to successfully restore intervertebral disc space and spinal alignment during the fusion process. The biomaterial should have high flexural strength, which is the ability to resist breakage during bending, and high compressive strength, which is the ability to resist compression under pressure, to withstand the static and dynamic forces exerted on the spine during daily activities over the long term.
- *Anti-Infective.* Spinal fusion devices can become colonized with bacteria, which may limit fusion to adjacent vertebrae or cause serious infection. Treating device-related infection is costly and generally requires repeat surgery, including surgery to replace the device, referred to as revision surgery, which may extend hospital stays, suffering and disability for patients. A biomaterial that has anti-infective properties can reduce the incidence of bacteria colonization in and around the interbody device that can lead to infection, revision surgery and associated increased costs. Publicly available articles report infection rates following implantation of traditional spinal fusion devices ranging from 3% to 18%.
- *Imaging Compatibility.* The biomaterial should be visible through, and not inhibit the effective use of, common surgical and diagnostic imaging techniques, such as x-ray, CT and MRI. These imaging techniques are used by surgeons during and after spinal fusion procedures to assist in the proper placement of interbody devices and to assess the quality of post-operative bone fusion.

Limitations of Biomaterials used in Interbody Spinal Fusion Devices

The three biomaterials most commonly used in interbody spinal fusion devices are PEEK, human cadaver bone, also referred to as allograft bone, and metals. We believe these materials do not possess the key characteristics required to form the optimal interbody spinal fusion device and are susceptible to potential fracture, implant-related infection, pain, limited fusion and instability, which have resulted in revision surgeries.

PEEK (polyetheretherketone)

PEEK is the most frequently used biomaterial for interbody spinal fusion devices and accounted for almost half of the devices implanted in the United States in 2012. We believe that the rate of revision surgery for PEEK interbody spinal fusion devices is approximately 6%. We believe this is caused by the following limitations of PEEK:

- *Restricts Bone Growth.* Due to PEEK's hydrophobic nature, the human body may recognize PEEK as a foreign substance and, therefore, may encapsulate the device with fibrous tissue. Although it is still possible for bone to grow through the device, bone may not adhere to the surface of the device if this tissue develops.
- *Lacks Strength and Resistance to Fracture.* PEEK lacks sufficient flexural strength, compressive strength and resistance to fracture necessary to reduce the risk of deformity or fracture during the fusion process. In addition, PEEK devices may fracture during implantation in certain interbody spinal fusion procedures. For example, in December 2012, Zimmer Spine recalled its PEEK Ardis® Interbody System Inserter, a surgical instrument used to implant a PEEK interbody spinal fusion device, because it resulted in the PEEK implants being susceptible to breakage when too much lateral force was applied to the inserter during implantation.
- *Lacks Anti-Infective Properties.* PEEK does not have any inherent anti-infective properties. In fact, a biofilm may form around a PEEK device that allows the colonization of bacteria, which can lead to infection.
- *Lacks Imaging Compatibility.* PEEK is invisible on x-rays. As a result, manufacturers of PEEK devices add metal markers to their devices so surgeons can see the location of the devices by x-ray. These markers, however, do not show the full outline of the device, which makes it difficult to assess the accuracy of the placement of the device. In addition, the metal markers cause artifacts on CT and MRI that can compromise the quality of the image.

Allograft Bone

Allograft bone is the second most frequently used biomaterial in interbody spinal fusion devices and accounted for over 40% of the devices implanted in the United States in 2012. Allograft bone has the following limitations:

- *Limited Promotion of Bone Growth.* Allograft bone has limited osteoinductive characteristics and therefore may not effectively promote bone growth in and around the interbody device.

- *Lacks Strength and Resistance to Fracture.* Generally, allograft bone is not as strong as live bone within the body or other materials used in interbody devices. In addition, techniques used to sterilize allograft bone, like gamma irradiation, can cause the allograft to become brittle and more likely to fracture.
- *Lacks Anti-Infective Properties and Risk of Disease Transmission.* In addition to not having inherent anti-infective properties, allograft bone exposes patients to a greater risk of disease transmission and auto-immune response.

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In addition, allograft bone is subject to inconsistent quality and size, which may require surgeons to make compromises on the fit of the device during surgery.

Metals

We believe metal interbody devices accounted for less than 10% of the devices implanted in the United States in 2012. Metals have the following limitations:

- *Limited Promotion of Bone Growth.* Metals have limited osteoinductive characteristics and therefore do not effectively promote bone growth in and around the interbody device.
- *Lack Anti-Infective Properties.* Metals do not have inherent anti-infective properties and do not suppress the colonization of bacteria in and around the device which can lead to infection.
- *Lack Imaging Compatibility.* Metals are opaque in x-rays and can cause significant imaging artifacts in CTs and MRIs. This can make it difficult for surgeons to detect the extent and quality of bone growth in and around the device in post-operative diagnostic imaging procedures.

The Hip and Knee Joint Replacement Market

Total joint replacement involves removing the diseased or damaged joint and replacing it with an artificial implant consisting of components made from several different types of biomaterials. The key components of a total hip implant include an artificial femoral head, consisting of a ball mounted on an artificial stem attached to the femur, and a liner, which is placed inside a cup affixed into the pelvic bone. The femoral head and liner move against each other to replicate natural motion in what is known as an articulating implant. Total knee replacement implants also use articulating components and are comprised of the following four main components: a femoral condyle, which is a specially shaped bearing that is affixed to the lower end of the femur; a tibial tray that is affixed to the upper end of the tibia; a tibial insert that is rigidly fixed to the tibial tray and serves as the surface against which the femoral condyle articulates; and a patella, or knee cap, which also articulates against the femoral condyle.

Implants for total hip and knee replacements are primarily differentiated by the biomaterials used in the components that articulate against one another. The combinations of biomaterials most commonly used in hip and knee replacement implants in the United States are metal-on-cross-linked polyethylene and traditional ceramic-on-cross-linked polyethylene. The use of hip replacement implants incorporating metal-on-metal and traditional ceramic-on-traditional ceramic biomaterials experienced a steep decline in the United States over the last several years due to their significant limitations. We believe that the most common currently used biomaterials in joint replacement implants also have limitations, and do not possess all of the following key characteristics required for optimal total joint replacement implants:

- *Resistance to Wear.* The biomaterials should have sufficient hardness and toughness, as well as extremely smooth surfaces, to effectively resist wear. Because the articulating implants move against each other, they are subject to friction, which frequently lead to abrasive wear and the release of small wear particles. This may cause an inflammatory response which results in osteolysis, or bone loss. Surgeons have identified osteolysis as a leading cause of joint implant failure, resulting in the need for revision surgery to replace the failed implant. One of the most commonly used combinations of biomaterials, metal-on-cross-linked polyethylene, as well as metal-on-metal implants tend to generate a large number of metal wear particles, which can cause osteolysis and a moderate to severe allergic reaction to the metal, referred to as metal sensitivity. While less common, metal implants may also cause a serious condition called metallosis. Both metal sensitivity and metallosis can result in revision surgery.
- *Non-Corrosive.* The biomaterials should be non-corrosive and should not cause adverse patient reactions. Metal placed in the human body corrodes over time and also results in the formation of metal ions, which leads to metal sensitivity in approximately 10% to 15% of the population and, less commonly, metallosis. As a result, there are significant increased risks from using metal-on-cross-linked polyethylene and metal-on-metal implants.
- *Hardness, Strength and Resistance to Fracture.* The biomaterials should be hard, strong and resistant to fracture to adequately bear the significant loads placed on joints like the hip and knee during daily activities. We believe there are strength limitations associated with traditional ceramic-on-cross-linked polyethylene and traditional ceramic-on-traditional ceramic implants.
- *Anti-Infective.* The biomaterials should have anti-infective properties to reduce the risk of bacteria colonization in and around the components that can lead to infection, revision surgeries and associated increased costs. Anti-infective properties reduce the

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risk of bacteria colonization in and around the components and reduce the likelihood of infection, revision surgeries and associated increased costs. None of the most commonly used biomaterials in joint replacement implants have anti-infective properties.

Our Silicon Nitride Technology Platform

We believe we are the first and only company to use silicon nitride in medical applications. Silicon nitride is a chemical compound comprised of the elements silicon and nitrogen, with the chemical formula Si_3N_4 . Silicon nitride, an advanced ceramic, is lightweight, resistant to fracture and strong, and is used in many demanding mechanical, thermal and wear applications, such as in space shuttle bearings, jet engine components and body armor.

We believe our silicon nitride is ideally suited for use in many medical applications and has the following characteristics that make it superior to other biomaterials, including PEEK, bone, metal and traditional ceramics, which do not possess all of these characteristics:

- *Promotes Bone Growth.* Our silicon nitride is osteoconductive through its inherent surface topography that provides support for new bone growth. We also believe our silicon nitride promotes an ideal environment for osteoinduction. As a hydrophilic material, silicon nitride attracts protein cells and nutrients that stimulate osteoprogenitor cells to differentiate into osteoblasts, which are needed for bone growth. Our silicon nitride also has an inherent surface chemistry that is more similar to bone than PEEK and metals are. As a result, we believe our silicon nitride has superior osteoconductive and osteoinductive properties when compared to other biomaterials, including those commonly used in interbody spinal fusion devices, such as PEEK, allograft bone and metal. These properties are highlighted in an *in vivo* study, where we measured the force required to separate devices from the spine after being implanted for three months, which indicates the level of osteointegration. In the absence of bacteria, the force required to separate our silicon nitride from its surrounding bone was approximately three times that of PEEK, and nearly two times that of titanium. In the presence of bacteria, the force required to separate our silicon nitride from its surrounding bone was over five times that of titanium, while there was effectively no separation force required for PEEK, indicating essentially no osteointegration.
- *Hard, Strong and Resistant to Fracture.* Our silicon nitride is hard, strong and possesses superior resistance to fracture over traditional ceramics and greater strength than polymers currently on the market. For example, our silicon nitride's flexural strength is more than five times that of PEEK and our silicon nitride's compressive strength is over twenty times that of PEEK. Unlike PEEK interbody spinal fusion devices, we believe our silicon nitride inbody spinal fusion devices can withstand the forces exerted during implantation and daily activities over the long term.
- *Anti-Infective.* We have demonstrated in *in vitro* and *in vivo* studies that silicon nitride has inherent anti-infective properties, which reduce the risk of infection in and around a silicon nitride device. PEEK, traditional ceramics, metals and bone do not have inherent anti-infective characteristics. These properties were highlighted in an *in vitro* study, where live bacteria counts were between 8 and 30 times lower on our silicon nitride than PEEK and up to 8 times lower on our silicon nitride than titanium. In addition to improving patient outcomes, we believe the anti-infective properties of our silicon nitride should make it an attractive biomaterial to hospitals and surgeons who are not reimbursed by third-party payors for the treatment of hospital-acquired infections. Additionally, silicon nitride is synthetic and, therefore, there is a lower risk of disease transmission through cross-contamination or of an adverse auto-immune response, sometimes associated with the use of allograft bone.
- *Imaging Compatible.* Our silicon nitride interbody spinal fusion devices are semi-radiolucent and clearly visible in x-rays, and produce no distortion under MRI and no scattering under CT. These characteristics enable an exact view of the device for precise intra-operative placement and post-operative bone fusion assessment in spinal fusion procedures. We believe these qualities provide surgeons with greater certainty of outcomes with our silicon nitride devices than with other biomaterials, such as PEEK and metals.
- *Resistant to Wear.* We believe our silicon nitride joint implant product candidates will have comparable or higher resistance to wear than metal-on-cross-linked polyethylene and traditional ceramic-on-cross-linked polyethylene joint implants, the two most commonly used total hip replacement implants. Also, debris associated with metal implants increases the risk of metal sensitivity and metallosis. Wear debris is a primary reason for early failures of metal and polymer articulating joint components.
- *Non-Corrosive.* Our silicon nitride does not have the drawbacks associated with the corrosive nature of metal within the body, including metal sensitivity and metallosis, nor does it result in the release of metal ions into the body. As a result, we believe our silicon nitride products will have lower revision rates and fewer complications than comparable metal products.

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Our Forms of Silicon Nitride

The chemical composition of our in-house formulation of silicon nitride, processing and manufacturing experience allow us to produce silicon nitride in four distinct forms. This capability provides us with the ability to utilize our silicon nitride in a variety of ways depending on its intended application, which, together with our silicon nitride's key characteristics, distinguishes us from manufacturers of products using other biomaterials.

We currently produce silicon nitride for use in our commercial products and product candidates in the following forms:

- *Solid Silicon Nitride, or MC²*. This form of silicon nitride is a fully dense, load-bearing solid, and is used for devices that require high strength, toughness, fracture resistance and low wear, including for interbody spinal fusion devices, hip and knee replacement implants and dental implants.



- *Porous Silicon Nitride, or C^SC*. While this form of silicon nitride has a chemical composition that is identical to that of MC², the C^SC form of silicon nitride has a porous structure, which is engineered to mimic cancellous bone, the spongy like bone tissue that typically makes up the interior of human bones. Our porous silicon nitride has interconnected pores ranging in size between about 90 and 600 microns, which is similar to that of cancellous bone. This form of silicon nitride can be used for the promotion of bone in-growth and attachment. Our porous silicon nitride is used as a substitute for the orthobiologics currently used to fill interbody devices in an effort to stimulate fusion and as a bone void filler, and as a porous scaffold for medical devices.



- *Composite Silicon Nitride*. This form of silicon nitride is a combination, or composite, of MC² and C^SC forms of silicon nitride. This composite may be used to manufacture devices and implants that mimic the structure of natural bone by incorporating both a fully dense, load-bearing solid MC² component on the outside and a porous C^SC component intended to promote bone in-growth on the inside. This composite form of silicon nitride is used in interbody spinal fusion devices and can be used in components for total hip and knee replacement implants.



- *Silicon Nitride Coating*. With a similar chemical composition as our other forms of silicon nitride, this form of silicon nitride can be applied as an adherent coating to metallic substrates, including cobalt-chromium, titanium and steel alloys. We believe applying silicon nitride as a coating may provide a highly wear-resistant articulation surface, such as on femoral heads, which may reduce problems associated with metal or polymer wear debris. We also believe that the silicon nitride coating can be applied to devices that require firm fixation and functional connections between the device or implant and the surrounding tissue, such as hip stems and screws. The use of silicon nitride coating may also create an anti-infective barrier between the device and the adjacent bone or tissue.



Our Competitive Strengths

We believe we can use our silicon nitride technology platform to become a leading biomaterial company and have the following principal strengths:

- *Sole Provider of Silicon Nitride Medical Devices.* We believe we are the only company that designs, develops, manufactures and sells medical grade silicon nitride-based products. Due to its key characteristics, we believe our silicon nitride enables us to offer new and transformative products across multiple medical specialties. In addition, with the FDA clearance of our silicon nitride *Valeo* products, we are one of only three companies that have developed and manufacture a ceramic for use in FDA cleared orthopedic medical devices in the United States.

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- *In-House Manufacturing Capabilities.* We operate a 30,000 square foot manufacturing facility located at our corporate headquarters in Salt Lake City, Utah. This operation complies with the FDA's quality system regulation, or QSR, and is certified under the International Organization for Standardization's, or ISO, standard 13485 for medical devices. This state-of-the-art facility allows us to rapidly design and produce silicon nitride products and control the entire manufacturing process from raw material to finished goods. We have also entered into a cooperative research and development agreement with Kyocera Industrial Ceramics Corporation, or Kyocera, under which we will work with Kyocera to determine its ability to become a second qualified manufacturer of our silicon nitride-based spinal fusion products and product candidates.
- *Established Commercial Infrastructure.* We market and sell our products to surgeons and hospitals in the United States, and select markets in Europe and South America through our established network of more than 50 independent sales distributors who are managed by our experienced in-house sales and marketing management team. As a result, our product revenue is driven by end-user prices, unlike other biomaterial companies that sell their products at lower prices to OEMs who then sell their products to the end user. Our control over the sales and marketing processes also allows us greater flexibility to selectively collaborate with distributors when we believe their experience or geographic reach can be beneficial to us.
- *Portfolio of Non-Silicon Nitride Products.* In addition to designing, developing, manufacturing and commercializing silicon nitride interbody spinal fusion devices, we sell a complementary line of non-silicon nitride spinal fusion products. We offer a full suite of spinal fusion products, which increases our access to surgeons and hospitals, and allows us to more effectively market our silicon nitride spinal fusion products to our customers. Product revenue from the sale of these non-silicon nitride products also supports further development of our silicon nitride products and product candidates.
- *Highly Experienced Management and Surgeon Advisory Team.* We have recently assembled a senior management team with over 150 years of collective experience in the healthcare industry. Members of our management team have experience in product development, launching new products into the orthopedics market and selling to hospitals through direct sales organizations, distributors, manufacturers and other orthopedic companies. We also collaborate with a network of leading surgeon advisors in the design and use of our products and product candidates.

Our Strategy

Our goal is to become a leading biomaterial company focused on using our silicon nitride technology platform to develop, manufacture and commercialize a broad range of medical devices. Key elements of our strategy to achieve this goal are the following:

- *Drive Further Adoption of our Silicon Nitride Interbody Spinal Fusion Devices.* We believe that increasing the awareness of our silicon nitride technology by educating surgeons about its key benefits, and design improvements to our silicon nitride products and related instruments, will accelerate the adoption of our products and ultimately help improve patient outcomes. We continue to innovate with further design enhancements in the introduction of our second-generation interbody spinal fusion devices. We are currently selling this new line of AL, PL and OL *Valeo* to select surgeons and expect to complete the full launch of the line in the United States in the second quarter of 2014. We are also completing the development of our second generation Cervical and TL *Valeo* interbody spinal fusion device and expect these to be launched in the second half of 2014. To drive further awareness of our products and the associated benefits offered by our silicon nitride technologies, we will continue to educate surgeons through multiple channels including industry conferences and meetings, media outlets and through our sales and marketing efforts. We also plan to facilitate the publication of data from bench testing and clinical outcome case studies.
- *Continue to Implement our Design and Build Program.* In the first half of 2013, we initiated a commercialization strategy, referred to as our Design and Build Program, in which we collaborate with influential surgeons to develop customized silicon nitride spinal fusion products and instruments. We first sell these products for use by the designing surgeons and a team of evaluating surgeons for their review based on their individual preferences focused on ease of use of the product and instrumentation and patient outcomes as compared to the previous products and instruments used by the surgeon. After the enhanced products are sold and evaluated and, if accepted by these surgeons, we plan to introduce these products more broadly into the market. The first products designed under this program were sold for initial evaluation in 2013.
- *Enhance our Commercial Infrastructure.* We expect to increase the productivity of our sales and marketing infrastructure to help us further penetrate the interbody spinal fusion market by continuing to engage experienced independent sales distributors with strong orthopedic surgeon relationships. For example, in October 2013, we entered into a new European sales agent agreement with K2M, Inc., one of the largest privately held spinal device companies in the world. We also periodically conduct programs to ensure that our distributors are knowledgeable about how the characteristics of our silicon nitride devices meet the demands of a range of spinal fusion procedures, and we regularly update our distributors about studies, test results, reviews and other developments that demonstrate the competitive advantages of our silicon nitride devices. We may also establish

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distribution collaborations in the United States and abroad when access to large or well-established sales and marketing organizations may help us gain access to new markets, increase sales in our existing markets, or accelerate market penetration for selected products.

- *Develop Silicon Nitride for Total Joint Components.* We are incorporating our silicon nitride technology into silicon nitride-coated components for use in total hip and knee replacement product candidates that we plan on developing in collaboration with a strategic partner. We also have designs for solid silicon nitride components and we will make a decision in the future about whether to pursue the development of these components. In December 2013, we participated in a pre-submission meeting with the FDA to finalize the regulatory strategy for a 510(k) clearance of our silicon nitride-coated total joint components in the United States. The FDA reviewers confirmed that the regulatory pathway would be a standard 510(k) clearance with supporting biomechanical testing. In response, we intend to develop silicon nitride-coated metal joint replacement components and then, together with a strategic partner, initiate biomechanical testing with our silicon nitride-coated metal components for use in total hip and knee replacement procedures to support a 510(k) submission to the FDA. We intend to pursue clearance of a total hip replacement product first, and if clearance is obtained, we intend to commercially launch products for use in total hip replacement by the second half of 2015.
- *Apply our Silicon Nitride Technology Platform to Other Opportunities.* Our silicon nitride technology platform is adaptable and we believe it may be used to develop products to address other significant opportunities, such as in the dental, sports medicine and trauma markets. We have manufactured prototypes of dental implants, sports medicine and trauma products, and we have developed a process to coat metals with our silicon nitride to enhance current medical devices and instruments. We plan to collaborate with other companies to develop and commercialize any future products in those areas or we may develop any one of them by ourselves if sufficient resources should become available.

Our Products and Product Candidates

We currently market a family of silicon nitride interbody spinal fusion devices and other non-silicon nitride spinal fusion products for use in cervical and lumbar spinal fusion surgical procedures to treat patients who suffer from degenerative, diseased and traumatic spine conditions. We are also developing multiple silicon nitride components for use in our total hip and knee replacement product candidates.

Spinal Fusion Products and Product Candidates

Our Valeo Silicon Nitride Products and Product Candidates

Our first generation *Valeo* silicon nitride spinal fusion device received 510(k) regulatory clearance and a CE mark in 2008. Based on surgeon feedback, we developed a second generation of *Valeo* products. In 2012, we received 510(k) clearance to market this second generation family of *Valeo* interbody spinal fusion devices, and we launched them with a select number of surgeons in 2013. Our second generation *Valeo* interbody spinal fusion devices offer distinct improvements over the first generation. The instrumentation of the second generation devices allow for better control of the device during implantation. The device allows for improved stability and potentially improved fusion after implantation and is offered in a broader selection of sizes. We expect to complete the full launch of our second generation AL, PL and OL *Valeo* interbody spinal fusion devices in the United States in the second quarter of 2014. We are also completing the development of our second generation Cervical and TL *Valeo* interbody spinal fusion devices and expect these to launch in the second half of 2014.

Our current products are:

<u>Valeo Interbody Fusion Devices</u>	<u>Generation</u>
AL: Anterior Lumbar	1st and 2nd
PL: Posterior Lumbar	1st and 2nd
OL: Oblique Lumbar	1st and 2nd
TL: Transforaminal Lumbar	1st and 2nd
C: Cervical	1st
CORP: Corpectomy	1st

We are also in the process of finishing the development of a *Valeo* stand-alone anterior lumbar intervertebral fusion device made from our *MC²* silicon nitride. The *Valeo* stand-alone product candidate, which incorporates fixation screws, will allow surgeons to perform less invasive procedures. We believe this may result in better patient outcomes compared to other spinal fusion procedures. We anticipate seeking 510(k) clearance for this product candidate in the second quarter of 2014, and, if cleared by the FDA, we anticipate launching our *Valeo* stand-alone product candidate in the United States in the second half of 2014.

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In 2009, we received a CE Mark to commercialize our *Valeo* interbody spinal fusion devices made from our composite silicon nitride. The porous *C^{SC}* center structure of these devices is designed to facilitate bone growth into the device, which we believe will allow surgeons to reduce or eliminate the use of allograft bone and other osteoconductive biomaterials. We are currently marketing these devices in the Netherlands, Spain and Germany. Additionally, we are conducting a prospective clinical trial in Europe, named CASCADE, comparing our *Valeo* composite silicon nitride interbody devices to PEEK interbody devices to obtain additional safety and efficacy data to support the 510(k) clearance in the United States. The trial is 100% enrolled. We expect results to be available in the second half of 2014. If this trial is successful, we plan to file a 510(k) submission with the FDA by mid-2015.



Valeo Composite (MC² + C^{2C})

Our Non-Silicon Nitride Spinal Fusion Products

We sell a line of complementary non-silicon nitride spinal fusion products to provide surgeons and hospitals with a broader range of products. Product revenue from the sale of our non-silicon nitride spinal fusion products further supports development of our silicon nitride products and product candidates. We plan to enhance our metal spinal fusion products with a silicon nitride coating. The following table lists our marketed non-silicon nitride spinal products.

<u>CATEGORY</u>	<u>PRODUCT NAME</u>	<u>BIOMATERIAL</u>
Facet Fixation System	Facet Gun Max/Facet Bolt Javelin: MIS Locking Facet System	Metal
Lumbar Spine Fixation	Preference Classic Spine System Preference 2 Spine System Preference 2 Complex Spine System	Metal
Orthobiologics	Preference Element Bone Graft Substitute BioDefense: Human Amnion Stem Cell Wound Covering Patch BioDlogics: Human Amnion Stem Cell Liquid Wound Covering <i>Valeo</i> BP: Synthetic Bone Putty PROCET: Facet Fusion Allograft Implant	Allograft
Interbody Spinal Fusion Device	Phantom Plus PLIF/TLIF IBFD Phantom Plus Cervical Spacer	PEEK

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Our Total Hip and Knee Joint Replacement Product Candidates

Our Total Hip Implant Product Candidates

We have developed two designs of femoral heads for use in our total hip replacement product candidates. Our first design is a silicon nitride-coated metal femoral head, for total joint replacement, which we plan to develop with a medical device partner. The second design is a femoral head that is made from our solid MC^2 silicon nitride and we are collaborating with Orthopaedic Synergy, Inc. to develop a total hip replacement product candidate using this design. These femoral heads are expected to articulate against a cross-linked polyethylene liner, fixed into a metal acetabular cup. We intend to initially advance our process to develop silicon nitride-coated femoral heads and then, together with a strategic partner, initiate biomechanical testing with our silicon nitride-coated femoral head for use in total hip replacement procedures to support a 510(k) submission to the FDA. If clearance is obtained, we intend to commercially launch products for use in total hip replacement by the second half of 2015. Although we have designs for solid silicon nitride components, we have not yet determined if we will pursue the development of these components.

Our Total Knee Implant Product Candidates

We have developed two designs of femoral condyle components for use in our total knee replacement product candidates. The first design utilizes our silicon nitride coating and we plan to partner with a medical device company to incorporate this design into a total knee replacement product candidate. The second design is made from our solid MC^2 silicon nitride and we are collaborating with Orthopaedic Synergy Inc. to develop a total knee joint replacement for this design. The femoral condyle component will attach to the lower end of the femur. The femoral condyle is expected to articulate against a cross-linked polyethylene tibial insert that will attach to the tibial tray at the upper end of the tibia, which we expect will be made from metal. We have successfully made prototypes of both designs. We intend to develop silicon nitride-coated femoral condyle components and then, together with a strategic partner, initiate biomechanical testing with our components for use in knee replacement procedures to support a 510(k) submission to the FDA. If clearance is obtained we intend to commercialize our products for use in total knee replacement surgeries post-FDA clearance. Although we have designs for solid silicon nitride components, we have not yet determined if we will pursue the development of these components.

Other Product Opportunities

Our silicon nitride technology platform is adaptable and we believe it may be used to develop products to address other significant opportunities, such as in the dental, sports medicine and trauma markets.

We also believe our coating technology may be used to enhance our marketed metal products as well as other commercially available metal spinal fusion and joint replacement products. We have produced feasibility prototypes of dental implants, other components for use in total hip implants in addition to our total hip and knee implant product candidates discussed above, a suture anchor for sports medicine and prototypes of silicon nitride-coated plates for potential trauma applications. We have also developed a process to apply our silicon nitride as a coating on other biomaterials.

The FDA has not evaluated any of these potential products and we are not currently advancing the development of any of these product candidates. We plan to collaborate with medical device companies to complete the development of and commercialize any product candidates we advance in these areas or develop any one of them ourselves if sufficient resources should become available.

Supporting Data

We and a number of independent third parties have conducted extensive biocompatibility, biomechanical, *in vivo* and *in vitro* testing on our silicon nitride to establish its safety and efficacy in support of regulatory clearance of our biomaterial, products and product candidates. We have also completed additional testing of our silicon nitride products and product candidates. The results of this testing have been published in peer review publications. Additionally, we have initiated prospective randomized clinical trials in humans *in vivo* and *in vitro* to support and expand our understanding of our silicon nitride's performance relative to other biomaterials and medical devices. We believe our product development strategy is consistent with the manner in which other biomaterials have been successfully introduced into the market and adopted as the standard of care. Listed below is an overview of some of the key testing completed on our silicon nitride biomaterial, products and product candidates to date, as well as other information about our silicon nitride and other biomaterials.

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Biocompatibility

Before our silicon nitride was first used in commercial products in 2008, we conducted a series of biocompatibility tests following the guidelines of the FDA and ISO and submitted the results to the FDA as part of the regulatory clearance process. These tests confirmed that our silicon nitride products meet required biocompatibility standards for human use.

Promotion of Bone Growth

In 2012, we conducted two separate studies at Brown University, the results of which suggest that the chemistry and inherent surface topography of our solid *MC²* silicon nitride provides an optimal environment for bone growth onto and around the device.

The first study was a series of *in vitro* analyses of protein adsorption, or presence on the surface of the biomaterial, onto silicon nitride, PEEK and titanium. The results of this study indicated that adsorption of two key proteins necessary for bone growth (fibronectin and vitronectin) were up to eight times greater on our silicon nitride than on PEEK, and up to four times greater than on titanium. A third important protein (laminin) had up to two times greater adsorption on our silicon nitride than on PEEK, and up to two-and-one-half times greater adsorption than on titanium.

The second study was an *in vivo* investigation of the osteointegration characteristics of these same three biomaterials after they had been surgically implanted into the skulls of laboratory rats. This study included an examination of the effect of *Staphylococcus epidermidis* bacteria on osteointegration. At time intervals of up to three months after implantation of the biomaterial, the amount of new bone growth within the surgical site and in direct contact with the implanted biomaterial was evaluated. In the absence of bacteria, new bone formation within the surgical site surrounding our silicon nitride was approximately 69%, compared with 36% and 24% for titanium and PEEK, respectively. Similarly, bone in direct contact, or apposition, with our silicon nitride, titanium and PEEK was 59%, 19% and 8%, respectively. As is common, in the presence of bacteria, new bone formation within the surgical site was suppressed, but still significantly greater for our silicon nitride than for the other two biomaterials. Observed new bone growth within the surgical site surrounding our silicon nitride was 41%, compared with 26% and 21% for titanium and PEEK, respectively. At the implant interface, the bone apposition for our silicon nitride, titanium and PEEK was 23%, 9% and 5%, respectively. To further characterize the extent of osteointegration, the force needed to separate each implant from its surrounding bone was measured. A larger force needed to separate the implant is an indication of improved osteointegration. At three months after implantation, in the absence of bacteria, the force required to separate our silicon nitride from its surrounding bone was approximately three times that of PEEK, and nearly two times that of titanium. In the presence of bacteria, there was effectively no separation force required for PEEK, indicating essentially no osteointegration. Our silicon nitride required over five times the force to separate it from its surrounding bone in the presence of bacteria in comparison to titanium.

In 2008, we conducted an animal study in which we evaluated the level of osteointegration of our porous *C^{SC}* silicon nitride with a knee-defect model in adult sheep. At three months after implantation, three out of five of the silicon nitride implants had extensive new bone formation at and into the implant surface, showing that the bone had grown into our *C^S C* silicon nitride to a depth of 3 millimeters, or mm. This animal study demonstrated the rapid osteointegration potential of our *C^{SC}* silicon nitride.

Hardness, Strength and Resistance to Fracture

Comparative Information

As shown in the table of comparative information publicly available about various biomaterials below:

- the hardness, or a material's resistance to deformity, of silicon nitride is comparable to traditional ceramics, but is substantially higher than either polymers or metals;
- the strength of silicon nitride is comparable or higher than metals and traditional ceramics, and is about sixteen to fifty-five times stronger than highly-cross-linked polyethylene, and four to eight times stronger than PEEK; and
- silicon nitride has the highest fracture resistance of any medical ceramic material and is three to eleven times more resistant to fracture than PEEK or highly-cross-linked polyethylene. This is due to the interwoven microstructure of silicon nitride. Metals have the highest fracture resistance.

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Comparison of Mechanical Properties Among Orthopedic Biomaterials

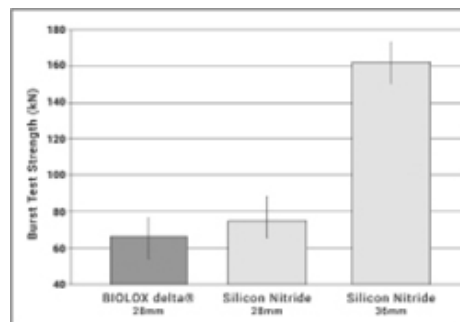
Material	Hardness (GPa)(1)	Strength (MPa)(1)	Fracture Resistance (MPa·m^{1/2})(1)
Silicon Nitride	13 – 16	800 – 1200	8 – 11
Aluminum Oxide Ceramic	14 – 19	300 – 500	3 – 5
Zirconia-Toughened Alumina Ceramic	12 – 19	700 – 1150	5 – 10
PEEK	0.09 – 0.28	160 – 180	2 – 3
Highly-Cross-Linked Polyethylene Polymer	0.03 – 0.07	22 – 48	1 – 2
Cobalt-Chromium Metal	3 – 4	700 – 1000	50 – 100
Titanium Alloy Metal	3 – 4	920 – 980	75

(1) GPa is a giga-pascal. MPa is a mega-pascal. Pascals are a measure of pressure. MPa·m^{1/2} is mega-pascal times a square root meter and is a measure related to the energy required to initiate fracture of a material.

We believe that the combination of high hardness, strength and fracture resistance positions our silicon nitride as an ideal biomaterial for many medical applications.

Burst Strength

In 2006, we conducted in-house comparative “burst strength” tests on femoral heads made from our silicon nitride produced by a contract manufacturer to our specifications and femoral heads made from one of the strongest commercially available ceramics, BIOLOX® *delta* (zirconia-toughened alumina). These tests were performed on three designs of 28 mm femoral heads using accepted testing protocols. The tests involved applying a load to each femoral head while mounted on a cobalt-chromium simulated hip implant stem, until the head burst. This enabled us to directly compare the strength of the femoral heads made of the two biomaterials. The results also provided an indication of each biomaterial’s resistance to fracture. The results of these tests are shown in the chart below.



The average burst test strength for the silicon nitride femoral heads in these tests was 75 kilonewtons, or kNs, compared with 65 kN for BIOLOX® *delta*, or about a 15% improvement. The burst strengths observed in our tests for BIOLOX® *delta* femoral heads are comparable to those observed by an independent party testing the same design BIOLOX® *delta* femoral heads as we did. We also conducted burst strength tests of 36 mm femoral heads made from our silicon nitride which showed those femoral heads had burst strengths that averaged 164 kN.

Resistance to Wear

In 2011, we commissioned an independent laboratory to conduct a wear study using our silicon nitride femoral heads. We tested our 28 mm silicon nitride femoral heads articulated against cross-linked polyethylene acetabular liners and our 40 mm silicon nitride femoral heads articulated against cross-linked polyethylene acetabular liners using well-established protocols in a hip simulator for their wear performance over 5 million cycles. We then compared the results for our silicon nitride product candidates to the results for the cobalt chrome femoral head and publicly available data from other commonly paired products. The results and comparison showed that:

- our silicon nitride-on-cross-linked polyethylene had approximately half the wear rate of that publicly reported for cobalt chrome-on-cross-linked polyethylene articulating hip components; and
- our silicon nitride-on-cross-linked polyethylene had comparable wear to that publicly reported for traditional ceramic-on-cross-linked polyethylene articulating hip components.

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Anti-Infective Properties

The results of the two studies at Brown University in 2012, demonstrate that our solid *MC*² silicon nitride has anti-infective properties. The objective of the *in vitro* study was to determine how our silicon nitride, PEEK and titanium interact with bacteria, protein and bone cells without the use of antibiotics and compared the growth of five different types of bacteria on silicon nitride, PEEK and titanium over time. Live bacteria counts were between 8 to 30 times lower on silicon nitride than PEEK and up to 8 times lower on silicon nitride than titanium.

In the *in vivo* study, bacteria were applied to the biomaterials before implantation. Three months after implantation, no infection was observed with silicon nitride, whereas both PEEK and titanium showed infection. The data demonstrate that our silicon nitride inhibits biofilm formation and bacterial colonization around the biomaterial.

Imaging Compatibility

In 2007, we conducted a study to compare the imaging characteristics of test blanks made of PEEK, the metals titanium and tantalum, and silicon nitride using a cadaver human vertebral body. Images of the vertebral body and the blanks were obtained using x-ray, CT and MRI under identical conditions. We assessed the radiolucent characteristics of the blanks in x-ray images quantitatively, assessed the presence of scatter in CT qualitatively and assessed distortion in MRI quantitatively. In x-ray, the metal blanks did not permit visualization of the underlying bone of the vertebral body, while PEEK was transparent, rendering its location difficult to determine. The silicon nitride blank had an intermediate radiolucency that rendered it visible and allowed a visual assessment of the underlying bone of the vertebral body. CT and MRI of the metal blanks indicated the presence of distortion while silicon nitride and PEEK exhibited no scattering.

Sales and Marketing

We market and sell our products to surgeons and hospitals through our established network of more than 50 independent sales distributors who are managed by our experienced 14 person in-house sales and marketing management team. Our sales efforts to date have been in the United States and selected markets in Europe and South America. To supplement our independent sales distributors, in select international markets, such as Europe, Japan, Australia and Canada, we may also seek to establish collaborations with leading orthopedic companies where we believe that a large, well-established partner may provide better access to those markets. For example, in October 2013, we entered into a European sales agent agreement with K2M, Inc., one of the largest privately held spinal device companies in the world. In addition, we may establish collaborations in the United States under circumstances where access to a larger sales and marketing organization may help to expand the market or accelerate penetration for selected products.

In the first quarter of 2013, we restructured the leadership of our sales and marketing team and hired a Senior Vice President of Global Sales, a Vice President of Marketing and a Senior Vice President, Strategic Marketing. This new leadership team has reviewed our entire sales and marketing practices and are implementing steps to improve the performance of these departments.

In addition to leveraging the strong existing surgeon relationships of our distribution network, we market our products through a combination of initiatives that are designed to establish and increase awareness of our silicon nitride products and their benefits over alternative products. We attend and make presentations at major industry events, including society meetings sponsored by the North American Spine Society, the America Academy of Orthopaedic Surgeons and the Congress of Neurological Surgeons, to educate surgeons and distributors about our products and product candidates. We advertise in trade journals and publications, and offer unique pricing strategies, including product bundling and incentivizing our distribution network to create and maintain long-term relationships with surgeons and hospitals. We also use surgeon advisors to assist in product development and to help implement awareness campaigns aimed at educating surgeons about our products. As part of these campaigns, we provide educational materials for hospitals and surgeons. We also conduct regional training seminars where our product managers, trainers, engineers, sales and marketing staff members work together with our surgeon advisors to educate surgeons and our distribution network in the use of our products.

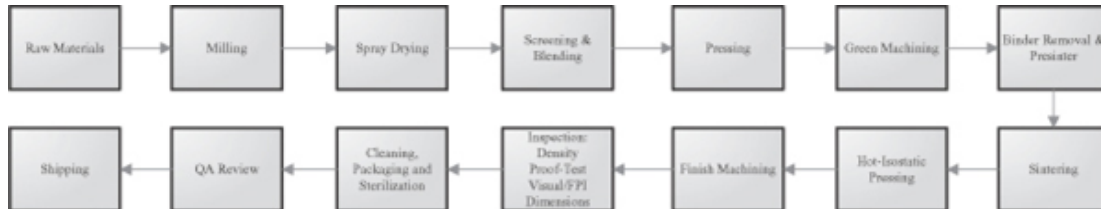
Manufacturing

Silicon Nitride Manufacturing

To control the quality, cost and availability of our silicon nitride products and product candidates, we operate our own manufacturing facility. Our 54,000 square foot corporate building includes a 30,000 square foot ISO 13485 certified medical device manufacturing space. It is equipped with state-of-the-art, powder processing, spray drying, pressing and computerized machining equipment, sintering furnaces, and other testing equipment that enables us to control the entire manufacturing process for our silicon

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nitride products and product candidates. To our knowledge, we are the only vertically integrated silicon nitride orthopedic medical device manufacturer in the world. All operations with the exceptions of raw material production, cleaning, packaging and sterilization are performed in-house. We purchase raw materials, consisting of silicon nitride ceramic powder and dopant chemical compounds, from several vendors which are ISO registered and approved by us. These raw materials are characterized and tested in our facility in accordance with our specifications and then blended to formulate our silicon nitride. We believe that there are multiple vendors that can supply us these raw materials and we continually monitor the quality and pricing offered by our vendors to ensure high quality and cost-effective supply of these materials. A flowchart of the silicon nitride manufacturing process is shown below.



In November 2013, we entered into a cooperative research and development agreement with Kyocera under which we will work with Kyocera to determine its ability to become a second qualified manufacturer of our silicon nitride-based spinal fusion products and product candidates.

Non-Silicon Nitride and Instruments Manufacturing

We obtain our non-silicon nitride spinal fusion products and instruments from third-party manufacturers. We also plan to rely on third-party manufacturers for the supply of the metal components of our silicon nitride hip and knee joint replacement product candidates. We only use manufacturers that operate under QSR and are ISO 13485 certified. Our in-house quality control group examines subcontracted components to ensure that they meet our required specifications. We believe that the use of third-party sources for non-silicon nitride spinal fusion products and instruments will reduce our capital investment requirements and allow us to strategically focus our resources on the manufacture of our silicon nitride products and product candidates.

Intellectual Property

We rely on a combination of patents, trademarks, trade secrets and other forms of intellectual property, nondisclosure agreements, proprietary information ownership agreements and other measures to protect our intellectual property rights. We believe that in order to have a competitive advantage, we must continue to develop and maintain the proprietary aspects of our technologies.

As of February 14, 2014, we had 34 issued U.S. patents, 38 pending U.S. patent applications, 11 granted foreign patents and 18 pending foreign patent applications. Our issued patents begin to expire in 2014, with the last of these patents expiring in 2031. The first core patents do not expire until 2022; these include US 6,881,229 and US 6,790,233.

We have seven U.S. patents, one European patent, and related pending applications, directed to articulating implants using our high-strength, high toughness doped silicon nitride MC^2 ceramic. The issued patents, which include US 6,881,229; US 7,666,229; US 8,123,812; US 7,780,738; US 7,695,521; US 7,776,085; US 8,133,284; and EP 1408874, begin to expire in 2022. We also have two U.S. patents, two European patents, and related pending applications, related to our C^SC technology that are directed to implants that have both a dense load-bearing, or cortical, component and a porous, or cancellous, component, together with a surface coating. The issued patents, which include US 6,790,233; US 6,846,327; EP 1389978; and EP 2055267, begin to expire in 2022.

We also have three U.S. patents that we acquired in July 2012 from Dytech Corporation Ltd., or Dytech, directed to manufacturing processes for the production of porous ceramics for use in our orthopedic implants. These patents, which include US 5,563,106; US 5,705,448; and US 6,617,270, expire between 2014 and 2019. Under our acquisition agreement with Dytech, Dytech granted to us a perpetual, irrevocable and exclusive license, including the right to grant sublicenses, to certain improvements and know-how related to the acquired patents. In return, we are required to pay Dytech a low single-digit royalty on net sales of products sold by us, our affiliates, or our licensees that are covered by one or more valid claims of these patents, and a percentage of any non-royalty licensing income we may receive in the event we grant a license to others.

Our remaining issued patents and pending applications are directed to additional aspects of our products and technologies including, among other things:

- designs for cervical plates;

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- designs for pedicle screws;
- designs for cervical disc implants;
- designs for intervertebral fusion devices;
- designs for facet fixation devices;
- designs for hip implants; and
- designs for knee implants.

We also expect to rely on trade secrets, know-how, continuing technological innovation and in-licensing opportunities to develop and maintain our intellectual property position. However, trade secrets are difficult to protect. We seek to protect the trade secrets in our proprietary technology and processes, in part, by entering into confidentiality agreements with commercial partners, collaborators, employees, consultants, scientific advisors and other contractors and into invention assignment agreements with our employees and some of our commercial partners and consultants. These agreements are designed to protect our proprietary information and, in the case of the invention assignment agreements, to grant us ownership of the technologies that are developed.

Competition

The main alternatives to our silicon nitride biomaterial include: PEEK, which is predominantly manufactured by Invibio, BIOLOX® *delta*, which is a traditional ceramic manufactured by CeramTec, allograft bone, metals and coated metals.

We believe our main competitors in the orthopedic implant market, which utilize a variety of competitive biomaterials, include: Medtronic, Inc.; DePuy Synthes Companies, a group of Johnson & Johnson companies; Stryker Corporation; Biomet, Inc.; Zimmer Holdings, Inc.; Smith & Nephew plc; and Aesculap Inc. Presently, these companies buy ceramic components on an OEM basis from manufacturers such as CeramTec, Kyocera and CoorTek, Inc., among others. We anticipate that these and other orthopedic companies and OEMs will seek to introduce new biomaterials and products that compete with ours.

Competition within the industry is primarily based on technology, innovation, product quality, and product awareness and acceptance by surgeons. Our principal competitors have substantially greater financial, technical and marketing resources, as well as significantly greater manufacturing capabilities than we do, and they may succeed in developing products that render our implants and product candidates non-competitive. Our ability to compete successfully will depend upon our ability to develop innovative products with advanced performance features based on our silicon nitride technologies.

Government Regulation of Medical Devices

Governmental authorities in the United States, at the federal, state and local levels, and other countries extensively regulate, among other things, the research, development, testing, manufacture, labeling, promotion, advertising, distribution, marketing and export and import of products such as those we are commercializing and developing. Failure to obtain approval or clearance to market our products and products under development and to meet the ongoing requirements of these regulatory authorities could prevent us from continuing to market or develop our products and product candidates.

United States

Pre-Marketing Regulation

In the United States, medical devices are regulated by the FDA. Unless an exemption applies, a new medical device will require either prior 510(k) clearance or approval of a premarket approval application, or PMA, before it can be marketed in the United States. The information that must be submitted to the FDA in order to obtain clearance or approval to market a new medical device varies depending on how the medical device is classified by the FDA. Medical devices are classified into one of three classes on the basis of the controls deemed by the FDA to be necessary to reasonably ensure their safety and effectiveness. Class I devices, which are those that have the lowest level or risk associated with them, are subject to general controls, including labeling, premarket notification and adherence to the QSR. Class II devices are subject to general controls and special controls, including performance standards. Class III devices, which have the highest level of risk associated with them, are subject to most of the previously identified requirements as well as to premarket approval. Most Class I devices and some Class II devices are exempt from the 510(k) requirement, although manufacturers of these devices are still subject to registration, listing, labeling and QSR requirements.

A 510(k) premarket notification must demonstrate that the device in question is substantially equivalent to another legally marketed device, or predicate device, that did not require premarket approval. In evaluating the 510(k), the FDA will determine whether the

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device has the same intended use as the predicate device, and (a) has the same technological characteristics as the predicate device, or (b) has different technological characteristics, and (i) the data supporting the substantial equivalence contains information, including appropriate clinical or scientific data, if deemed necessary by the FDA, that demonstrates that the device is as safe and as effective as a legally marketed device, and (ii) does not raise different questions of safety and effectiveness than the predicate device. Most 510(k)s do not require clinical data for clearance, but the FDA may request such data. The FDA's goal is to review and act on each 510(k) within 90 days of submission, but it may take longer based on requests for additional information. In addition, requests for additional data, including clinical data, will increase the time necessary to review the notice. If the FDA does not agree that the new device is substantially equivalent to the predicate device, the new device will be classified in Class III, and the manufacturer must submit a PMA. Since July 2012, however, with the enactment of the Food and Drug Administration Safety and Innovation Act, or FDASIA, a *de novo* pathway is directly available for certain low to moderate risk devices that do not qualify for the 510(k) pathway due to lack of a predicate device. Modifications to a 510(k)-cleared medical device may require the submission of another 510(k) or a PMA if the changes could significantly affect the safety or effectiveness or constitute a major change in the intended use of the device.

Modifications to a 510(k)-cleared device frequently require the submission of a traditional 510(k), but modifications meeting certain conditions may be candidates for FDA review under a Special 510(k). If a device modification requires the submission of a 510(k), but the modification does not affect the intended use of the device or alter the fundamental scientific technology of the device, then summary information that results from the design control process associated with the cleared device can serve as the basis for clearing the application. A Special 510(k) allows a manufacturer to declare conformance to design controls without providing new data. When the modification involves a change in material, the nature of the "new" material will determine whether a traditional or Special 510(k) is necessary. For example, in its Device Advice on How to Prepare a Special 510(k), the FDA uses the example of a change in a material in a finger joint prosthesis from a known metal alloy to a ceramic that has not been used in a legally marketed predicate device as a type of change that should not be submitted as a Special 510(k). However, if the "new" material is a type that has been used in other legally marketed devices within the same classification for the same intended use, a Special 510(k) is appropriate. The FDA gives as an example a manufacturer of a hip implant who changes from one alloy to another that has been used in another legally marketed predicate. Special 510(k)s are typically processed within 30 days of receipt.

The PMA process is more complex, costly and time consuming than the 510(k) clearance procedure. A PMA must be supported by extensive data including, but not limited to, technical, preclinical, clinical, manufacturing, control and labeling information to demonstrate to the FDA's satisfaction the safety and effectiveness of the device for its intended use. After a PMA is submitted, the FDA has 45 days to determine whether it is sufficiently complete to permit a substantive review. If the PMA is complete, the FDA will file the PMA. The FDA is subject to performance goal review times for PMAs and may issue a decision letter as a first action on a PMA within 180 days of filing, but if it has questions, it will likely issue a first major deficiency letter within 150 days of filing. It may also refer the PMA to an FDA advisory panel for additional review, and will conduct a preapproval inspection of the manufacturing facility to ensure compliance with the QSR, either of which could extend the 180-day response target. While the FDA's ability to meet its performance goals has generally improved during the past few years, it may not meet these goals in the future. A PMA can take several years to complete and there is no assurance that any submitted PMA will ever be approved. Even when approved, the FDA may limit the indication for which the medical device may be marketed or to whom it may be sold. In addition, the FDA may request additional information or request the performance of additional clinical trials before it will reconsider the approval of the PMA or as a condition of approval, in which case the trials must be completed after the PMA is approved. Changes to the device, including changes to its manufacturing process, may require the approval of a supplemental PMA.

If a medical device is determined to present a "significant risk," the manufacturer may not begin a clinical trial until it submits an investigational device exemption, or IDE, to the FDA and obtains approval of the IDE from the FDA. The IDE must be supported by appropriate data, such as animal and laboratory testing results and include a proposed clinical protocol. These clinical trials are also subject to the review, approval and oversight of an institutional review board, or IRB, which is an independent and multi-disciplinary committee of volunteers who review and approve research proposals, and the reporting of adverse events and experiences, at each institution at which the clinical trial will be performed. The clinical trials must be conducted in accordance with applicable regulations, including but not limited to the FDA's IDE regulations and current good clinical practices. A clinical trial may be suspended by the FDA, the IRB or the sponsor at any time for various reasons, including a belief that the risks to the study participants outweigh the benefits of participation in the trial. Even if a clinical trial is completed, the results may not demonstrate the safety and efficacy of a device, or may be equivocal or otherwise not be sufficient to obtain approval.

Post-Marketing Regulation

After a device is placed on the market, numerous regulatory requirements apply. These include:

- compliance with the QSR, which require manufacturers to follow stringent design, testing, control, documentation, record maintenance, including maintenance of complaint and related investigation files, and other quality assurance controls during the manufacturing process;

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- labeling regulations, which prohibit the promotion of products for uncleared or unapproved or “off-label” uses and impose other restrictions on labeling; and
- medical device reporting obligations, which require that manufacturers investigate and report to the FDA adverse events, including deaths, or serious injuries that may have been or were caused by a medical device and malfunctions in the device that would likely cause or contribute to a death or serious injury if it were to recur.

Failure to comply with applicable regulatory requirements can result in enforcement action by the FDA, which may include any of the following sanctions:

- warning letters;
- fines, injunctions, and civil penalties;
- recall or seizure of our products;
- operating restrictions, partial suspension or total shutdown of production;
- refusal to grant 510(k) clearance or PMA approvals of new products;
- withdrawal of 510(k) clearance or PMA approvals; and
- criminal prosecution.

To ensure compliance with regulatory requirements, medical device manufacturers are subject to market surveillance and periodic, pre-scheduled and unannounced inspections by the FDA, and these inspections may include the manufacturing facilities of our subcontractors.

International Regulation

International sales of medical devices are subject to foreign government regulations, which vary substantially from country to country. The time required to obtain approval by a foreign country may be longer or shorter than that required for FDA approval, and the requirements may differ. For example, the primary regulatory authority with respect to medical devices in Europe is that of the European Union. The European Union consists of 28 countries and has a total population of over 500 million people. The unification of these countries into a common market has resulted in the unification of laws, standards and procedures across these countries, which may expedite the introduction of medical devices like those we are offering and developing. Norway, Iceland, Lichtenstein and Switzerland are not members of the European Union, but have transposed applicable European medical device laws into their national legislation. Thus, a device that is marketed in the European Union may also be recognized and accepted in those four non-member European countries as well.

The European Union has adopted numerous directives and standards regulating the design, manufacture, clinical trials, labeling and adverse event reporting for medical devices. Devices that comply with the requirements of relevant directives will be entitled to bear CE Conformity Marking, indicating that the device conforms to the essential requirements of the applicable directives and, accordingly, can be commercially distributed throughout the European Union. Actual implementation of these directives, however, may vary on a country-by-country basis. The CE Mark is a mandatory conformity mark on medical devices distributed and sold in the European Union and certifies that a medical device has met applicable requirements.

The method of assessing conformity varies, but normally involves a combination of self-assessment by the manufacturer and a third-party assessment by a “Notified Body.” Notified Bodies are independent testing houses, laboratories, or product certifiers authorized by the European Union member states to perform the required conformity assessment tasks, such as quality system audits and device compliance testing. An assessment by a Notified Body based within the European Union is required in order for a manufacturer to distribute the product commercially throughout the European Union. Medium and higher risk devices require the intervention of a Notified Body which will be responsible for auditing the manufacturer’s quality system. The Notified Body will also determine whether or not the product conforms to the requirements of the applicable directives. Devices that meet the applicable requirements of E.U. law and have undergone the appropriate conformity assessment routes will be granted CE “certification.”

The CE Mark is mandatory for medical devices sold not only within the countries of the European Union but more generally within most of Europe. As many of the European standards are converging with international standards, the CE Mark is often used on medical devices manufactured and sold outside of Europe (notably in Asia that exports many manufactured products to Europe). CE Marking gives companies easier access into not only the European market but also to Asian and Latin American markets, most of whom recognize the CE Mark on medical device as a mark of quality and adhering to international standards of consumer safety, health or environmental requirements.

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In September 2012, the European Commission adopted a proposal for a regulation which, if adopted, will change the way that most medical devices are regulated in the European Union, and may subject our products to additional requirements.

Compliance with Healthcare Laws

We must comply with various U.S. federal and state laws, rules and regulations pertaining to healthcare fraud and abuse, including anti-kickback and false claims laws, rules, and regulations, as well as other healthcare laws in connection with the commercialization of our products. Fraud and abuse laws are interpreted broadly and enforced aggressively by various state and federal agencies, including the U.S. Department of Justice, the U.S. Office of Inspector General for the Department of Health and Human Services and various state agencies.

We have entered into agreements with certain surgeons for assistance with the design of our products, some of whom we anticipate may make referrals to us or order our products. A majority of these agreements contain provisions for the payments of royalties and/or stock options. In addition, some surgeons currently own shares of our stock. We have structured these transactions with the intention of complying with all applicable laws, including fraud and abuse, data privacy and security, and transparency laws. Despite this intention, there can be no assurance that a particular government agency or court would determine our practices to be in full compliance with such laws. We could be materially impacted if regulatory or enforcement agencies or courts interpret our financial arrangements with surgeons to be in violation of healthcare laws, including, without limitation, fraud and abuse, data privacy and security, or transparency laws.

The U.S. federal Anti-Kickback Statute prohibits persons, including a medical device manufacturer (or a party acting on its behalf), from knowingly or willfully soliciting, receiving, offering or paying remuneration, directly or indirectly, in exchange for or to induce either the referral of an individual for a service or product or the purchasing, ordering, arranging for, or recommending the ordering of, any service or product for which payment may be made by Medicare, Medicaid or any other federal healthcare program. This statute has been interpreted to apply to arrangements between medical device manufacturers on one hand and healthcare providers on the other. The term "remuneration" is not defined in the federal Anti-Kickback Statute and has been broadly interpreted to include anything of value, such as cash payments, gifts or gift certificates, discounts, waiver of payments, credit arrangements, ownership interests, the furnishing of services, supplies or equipment, and the provision of anything at less than its fair market value. Courts have broadly interpreted the scope of the law, holding that it may be violated if merely "one purpose" of an arrangement is to induce referrals, irrespective of the existence of other legitimate purposes. The Anti-Kickback Statute prohibits many arrangements and practices that are lawful in businesses outside of the healthcare industry. Although there are a number of statutory exemptions and regulatory safe harbors protecting certain business arrangements from prosecution, the exemptions and safe harbors are drawn narrowly, and practices that involve remuneration intended to induce prescribing, purchasing or recommending may be subject to scrutiny if they do not qualify for an exemption or safe harbor. Our practices may not in all cases meet all of the criteria for safe harbor protection from federal Anti-Kickback Statute liability. The reach of the Anti-Kickback Statute was broadened by the recently enacted Patient Protection and Affordable Care Act of 2010 and the Health Care and Education Affordability Reconciliation Act of 2010, collectively, the Affordable Care Act or ACA, which, among other things, amends the intent requirement of the federal Anti-Kickback Statute such that a person or entity no longer needs to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation. In addition, the ACA provides that the government may assert that a claim including items or services resulting from a violation of the federal Anti-Kickback Statute constitutes a false or fraudulent claim for purposes of the federal False Claims Act (discussed below) or the civil monetary penalties statute, which imposes fines against any person who is determined to have presented or caused to be presented claims to a federal healthcare program that the person knows or should know is for an item or service that was not provided as claimed or is false or fraudulent. In addition to the federal Anti-Kickback Statute, many states have their own anti-kickback laws. Often, these laws closely follow the language of the federal law, although they do not always have the same scope, exceptions, safe harbors or sanctions. In some states, these anti-kickback laws apply not only to payments made by government healthcare programs but also to payments made by other third-party payors, including commercial insurance companies.

Sales, marketing, consulting, and advisory arrangements between medical device manufacturers and sales agents and physicians are subject to the Anti-Kickback Statute and other fraud and abuse laws. Government officials have focused recent enforcement efforts on, among other things, the sales and marketing activities of healthcare companies, including medical device manufacturers, and have brought cases against individuals or entities whose personnel allegedly offered unlawful inducements to potential or existing customers in an attempt to procure their business. We expect these activities to continue to be a focus of government enforcement efforts. Settlements of these cases by healthcare companies have involved significant fines and penalties and in some instances criminal plea agreements. We are also aware of governmental investigations of some of the largest orthopedic device companies reportedly focusing on consulting and service agreements between these companies and orthopedic surgeons. These developments are ongoing and we cannot predict the effects they will have on our business.

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The federal False Claims Act imposes liability on any person that, among other things, knowingly presents, or causes to be presented, a false or fraudulent claim for payment by a federal healthcare program. The *qui tam* provisions of the False Claims Act allow a private individual to bring civil actions on behalf of the federal government alleging that the defendant has submitted a false claim, or has caused such a claim to be submitted, to the federal government, and to share in any monetary recovery. There are many potential bases for liability under the False Claims Act. Liability arises, primarily, when a person knowingly submits, or causes another to submit, a false claim for reimbursement to the federal government. The False Claims Act has been used to assert liability on the basis of inadequate care, kickbacks, and other improper referrals, and allegations as to misrepresentations with respect to the services rendered. *Qui tam* actions have increased significantly in recent years, causing greater numbers of healthcare companies, including medical device manufacturers, to defend false claim actions, pay damages and penalties, or be excluded from participation in Medicare, Medicaid or other federal or state healthcare programs as a result of investigations arising out of such actions. In addition, various states have enacted similar laws analogous to the False Claims Act. Many of these state laws apply where a claim is submitted to any third-party payor and not merely a federal healthcare program. We are unable to predict whether we would be subject to actions under the False Claims Act or a similar state law, or the impact of such actions. However, the cost of defending such claims, as well as any sanctions imposed, could adversely affect our financial performance. The Health Insurance Portability and Accountability Act of 1996, or HIPAA, also created several new federal crimes, including healthcare fraud and false statements relating to healthcare matters. The healthcare fraud statute prohibits knowingly and willfully executing a scheme to defraud any healthcare benefit program, including private third party payors. The false statements statute prohibits knowingly and willfully falsifying, concealing, or covering up a material fact or making any materially false, fictitious, or fraudulent statement in connection with the delivery of or payment for healthcare benefits, items, or services.

In addition, we may be subject to, or our marketing or research activities may be limited by, data privacy and security regulation by both the federal government and the states in which we conduct our business. For example, HIPAA and its implementing regulations established uniform federal standards for certain “covered entities” (healthcare providers, health plans and healthcare clearinghouses) governing the conduct of certain electronic healthcare transactions and protecting the security and privacy of protected health information. The American Recovery and Reinvestment Act of 2009, commonly referred to as the economic stimulus package, included expansion of HIPAA’s privacy and security standards called the Health Information Technology for Economic and Clinical Health Act, or HITECH, which became effective on February 17, 2010. Among other things, HITECH makes HIPAA’s privacy and security standards directly applicable to “business associates”—independent contractors or agents of covered entities that create, receive, maintain, or transmit protected health information in connection with providing a service for or on behalf of a covered entity. HITECH also increased the civil and criminal penalties that may be imposed against covered entities, business associates and possibly other persons, and gave state attorneys general new authority to file civil actions for damages or injunctions in federal courts to enforce the federal HIPAA laws and seek attorney’s fees and costs associated with pursuing federal civil actions. These laws also require the reporting of breaches of protected health information to affected individuals, regulators and in some cases, local or national media. HIPAA and HITECH impose strict limits on our physician collaborators’ ability to use and disclose patient information on our behalf.

There are also an increasing number of state “sunshine” laws that require manufacturers to provide reports to state governments on pricing and marketing information. Several states have enacted legislation requiring medical device companies to, among other things, establish marketing compliance programs, file periodic reports with the state, make periodic public disclosures on sales and marketing activities, and to prohibit or limit certain other sales and marketing practices. In addition, a federal law known as the Physician Payments Sunshine Act, now requires medical device manufacturers to track and report to the federal government certain payments and other transfers of value made to physicians and teaching hospitals and ownership or investment interests held by physicians and their immediate family members. The first reporting period will cover only payments or transfers of value made and ownership or investment interests held by physicians and their immediate family members from August 1, 2013 to December 31, 2013. The federal government will disclose the reported information on a publicly available website beginning in September 2014. For calendar year 2014, the Physician Payments Sunshine Act will require medical device manufacturers to report payments and transfers of values made and ownership or investment interests held by physicians and their immediate family members for the full calendar year. These laws may adversely affect our sales, marketing, and other activities by imposing administrative and compliance burdens on us. If we fail to track and report as required by these laws or to otherwise comply with these laws, we could be subject to the penalty provisions of the pertinent state and federal authorities.

Clinical research is heavily regulated by FDA regulations for the protection of human subjects (21 C.F.R. 50 and 56) and also the regulations of the U.S Department of Health and Human Services, or the Common Rule (45 C.F.R 46). Both FDA human subject regulations and the Common Rule impose restrictions on the involvement of human subjects in clinical research and require, among other things, the balancing of the risks and benefits of research, the documented informed consent of research participants, initial and ongoing review of research by an IRB. Similar regulations govern research conducted in foreign countries. Compliance with human subject protection regulations is costly and time consuming. Failure to comply could substantially and adversely impact our research program and the development of our products.

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Because of the breadth of these laws and the narrowness of available statutory and regulatory exemptions, it is possible that some of our business activities could be subject to challenge under one or more of such laws. If our operations are found to be in violation of any of the federal and state laws described above or any other governmental regulations that apply to us, we may be subject to penalties, including criminal and significant civil monetary penalties, damages, fines, imprisonment, exclusion from participation in government healthcare programs, injunctions, recall or seizure of products, total or partial suspension of production, denial or withdrawal of pre-marketing product clearances and approvals, private “qui tam” actions brought by individual whistleblowers in the name of the government or refusal to allow us to enter into supply contracts, including government contracts, and the curtailment or restructuring of our operations. Public disclosure of privacy and data security violations could cause significant reputational harm. Any of these events could adversely affect our ability to operate our business and our results of operations. To the extent that any of our products are sold in a foreign country, we may be subject to similar foreign laws and regulations, which may include, for instance, applicable post-marketing requirements, including safety surveillance, anti-fraud and abuse laws, implementation of corporate compliance programs, as well as laws and regulations requiring transparency of pricing and marketing information and governing the privacy and security of health information, such as the E.U.’s Directive 95/46 on the Protection of Individuals with regard to the Processing of Personal Data, or the Data Directive, and the wide variety of national laws implementing the Data Directive.

Healthcare Reform

In the United States and foreign jurisdictions, there have been a number of legislative and regulatory changes to the healthcare system that could affect our future results of operations. In particular, there have been and continue to be a number of initiatives at the U.S. federal and state levels that seek to reduce healthcare costs.

In March 2010, President Obama signed into law the ACA, a sweeping law intended to broaden access to health insurance, reduce or constrain the growth of healthcare spending, enhance remedies against healthcare fraud and abuse, add new transparency requirements for healthcare and health insurance industries, impose new taxes and fees on pharmaceutical and medical device manufacturers and impose additional health policy reforms. Among other things, the ACA imposes a 2.3% medical device excise tax on sales of many medical devices in the United States which became effective on January 1, 2013. Substantial new provisions affecting compliance have also been enacted, which may affect our business practices with healthcare practitioners and a significant number of provisions are not yet, or have only recently become, effective. Although it is too early to determine the full effect of the ACA, the new law appears likely to place downward pressure on pricing of medical devices, especially under the Medicare program, and may also increase our regulatory burdens and operating costs.

In addition, other legislative changes have been proposed and adopted since the ACA was enacted. For example, on August 2, 2011, the President signed into law the Budget Control Act of 2011, which, among other things, created the Joint Select Committee on Deficit Reduction to recommend to Congress proposals in spending reductions. The Joint Select Committee on Deficit Reduction did not achieve a targeted deficit reduction of at least \$1.2 trillion for the years 2013 through 2021, triggering the legislation’s automatic reduction to several government programs. This includes aggregate reductions to Medicare payments to providers of up to 2% per fiscal year, starting in 2013. On January 2, 2013, President Obama signed into law the American Taxpayer Relief Act of 2012, or ATRA, which, among other things, reduced Medicare payments to several types of providers and increased the statute of limitations period for the government to recover overpayments to providers from three to five years. On March 1, 2013, the President signed an executive order implementing the Budget Control Act’s 2% Medicare payment reductions, and on April 1, 2013, these reductions went into effect. These new laws may result in additional reductions in Medicare and other healthcare funding, which could have a material adverse effect on our financial operations.

We expect that the ACA, as well as other healthcare reform measures that have been and may be adopted in the future, may result in more rigorous coverage criteria and in additional downward pressure on the price that we receive for our products. Any reduction in reimbursement from Medicare or other government programs may result in a similar reduction in payments from private payors. The implementation of cost containment measures or other healthcare reforms may affect our ability to generate revenue and profits or commercialize our product candidates.

Third-Party Reimbursement

Because we typically receive payment directly from hospitals and surgical centers, we do not anticipate relying directly on payment for any of our products from third-party payors, such as Medicare, Medicaid, private insurers, and managed care companies. However, our business will be affected by policies administered by federal and state healthcare programs, such as Medicare and Medicaid, as well as private third-party payors, which often follow the policies of the state and federal healthcare programs. For example, our business will be indirectly impacted by the ability of a hospital or medical facility to obtain coverage and third-party reimbursement for procedures performed using our products. Many hospitals and clinics in the United States belong to group purchasing organizations (that typically incentivize their hospital members to make a relatively large proportion of purchases from a limited

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number of vendors of similar products that have contracted to offer discounted prices). Such contracts often include exceptions for purchasing certain innovative new technologies, however. Accordingly, the commercial success of our products may also depend to some extent on our ability to either negotiate favorable purchase contracts with key group purchasing organizations or persuade hospitals and clinics to purchase our product “off contract.” These third-party payors may deny reimbursement if they determine that a device used in a procedure was not medically necessary; was not used in accordance with cost-effective treatment methods, as determined by the third-party payor; or was used for an unapproved use. A national or local coverage decision denying Medicare coverage for one or more of our products could result in private insurers and other third party payors also denying coverage. Even if favorable coverage and reimbursement status is attained for our products, less favorable coverage policies and reimbursement rates may be implemented in the future. The cost containment measures that third-party payors and providers are instituting, both within the United States and abroad, could significantly reduce our potential revenues from the sale of our products and any product candidates. We cannot provide any assurances that we will be able to obtain and maintain third party coverage or adequate reimbursement for our products and product candidates in whole or in part.

For inpatient and outpatient procedures, including those that will involve use of our products, Medicare and many other third-party payors in the United States reimburse hospitals at a prospectively determined amount. This amount is generally based on one or more diagnosis related groups, or DRGs, associated with the patient’s condition for inpatient treatment and generally based on ambulatory payment classifications, or APCs, associated with the procedures performed as an outpatient at an ambulation surgicenter. Each DRG or APC is associated with a level of payment and may be adjusted from time to time, usually annually. Prospective payments are intended to cover most of the non-physician hospital costs incurred in connection with the applicable diagnosis and related procedures. Implant products, such as those we plan to sell, represent part of the total procedure costs while labor, hospital room and board, and other supplies and services represent the balance of those costs. However, the prospective payment amounts are typically set independently of a particular hospital’s actual costs associated with treating a particular patient and implanting a device. Therefore, the payment that a hospital would receive for a particular hospital visit would not typically take into account the cost of our products.

Medicare has established a number of DRGs for inpatient procedures that involve the use of products similar to ours. Although Medicare has authority to create special DRGs for hospital services that more properly reflect the actual costs of expensive or new-technology devices implanted as part of a procedure, it has declined to do so in the past, and we do not expect that it will do so with respect to our current products and product candidates. Medicare’s DRG and APC classifications may have implications outside of Medicare, as many other U.S. third-party payors often use Medicare DRGs and APCs for purposes of determining reimbursement.

We believe that orthopedic implants generally have been well received by third-party payors because of the ability of these implants to greatly reduce long-term healthcare costs for patients with degenerative joint disease. However, coverage and reimbursement policies vary from payor to payor and are subject to change. As discussed above, hospitals that purchase medical devices for treatment of their patients generally rely on third-party payors to reimburse all or part of the costs and fees associated with the procedures performed with these devices. Both government and private third-party coverage and reimbursement levels are critical to new product acceptance. Neither hospitals nor surgeons are likely to use our products if they do not receive reimbursement for the procedures adequate to cover the cost of our products.

While it is expected that hospitals will be able to obtain coverage for procedures using our products, the level of payment available to them for such procedures may change over time. State and federal healthcare programs, such as Medicare and Medicaid, closely regulate provider payment levels and have sought to contain, and sometimes reduce, payment levels. Commercial insurers and managed care plans frequently follow government payment policies, and are likewise interested in controlling increases in the cost of medical care. These third-party payors may deny payment if they determine that a procedure was not medically necessary, a device used in a procedure was not used in accordance with cost-effective treatment methods, as determined by the third-party payor, or was used for an unapproved use.

In addition, some payors are adopting pay-for-performance programs that differentiate payments to healthcare providers based on the achievement of documented quality-of-care metrics, cost efficiencies, or patient outcomes. These programs are intended to provide incentives to providers to find ways to deliver the same or better results while consuming fewer resources. As a result of these programs, and related payor efforts to reduce payment levels, hospitals and other providers are seeking ways to reduce their costs, including the amounts they pay to medical device suppliers. Adverse changes in payment rates by payors to hospitals could adversely impact our ability to market and sell our products and negatively affect our financial performance.

In international markets, healthcare payment systems vary significantly by country and many countries have instituted price ceilings on specific product lines. There can be no assurance that our products will be considered cost-effective by third-party payors, that reimbursement will be available or, if available, that the third-party payors’ reimbursement policies will not adversely affect our ability to sell our products profitably.

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Member countries of the European Union offer various combinations of centrally financed healthcare systems and private health insurance systems. The relative importance of government and private systems varies from country to country. Governments may influence the price of medical devices through their pricing and reimbursement rules and control of national healthcare systems that fund a large part of the cost of those products to consumers. Some jurisdictions operate positive and negative list systems under which products may be marketed only once a reimbursement price has been agreed upon. Some of these countries may require, as condition of obtaining reimbursement or pricing approval, the completion of clinical trials that compare the cost-effectiveness of a particular product candidate to currently available therapies. Some E.U. member states allow companies to fix their own prices for devices, but monitor and control company profits. The choice of devices is subject to constraints imposed by the availability of funds within the purchasing institution. Medical devices are most commonly sold to hospitals or healthcare facilities at a price set by negotiation between the buyer and the seller. A contract to purchase products may result from an individual initiative or as a result of a competitive bidding process. In either case, the purchaser pays the supplier, and payment terms vary widely throughout the European Union. Failure to obtain favorable negotiated prices with hospitals or healthcare facilities could adversely affect sales of our products.

Employees

As of January 15, 2014, we had 79 employees, including five part-time temporary employees, of which 15 are employed in administration, 18 in operations, 35 in manufacturing and research and development, and 11 in sales and marketing. We believe that our success will depend, in part, on our ability to attract and retain qualified personnel. We have never experienced a work stoppage due to labor difficulties and believe that our relations with our employees are good. None of our employees are represented by labor unions.

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ITEM 1A. RISK FACTORS

In addition to the other information contained in this Annual Report, the following risk factors should be considered carefully in evaluating our company. Our business, financial condition, liquidity or results of operations could be materially adversely affected by any of these risks.

Risks Related to Our Business and Strategy

We have incurred net losses since our inception and anticipate that we will continue to incur substantial net losses for the foreseeable future. We may never achieve or sustain profitability.

We have incurred substantial net losses since our inception. For the years ended December 31, 2013 and 2012 we incurred a net loss of \$8.2 million and \$35.0 million, respectively, and used cash in operations of \$9.2 million and \$9.7 million, respectively. We have an accumulated deficit of \$139.9 million at December 31, 2013. With the exception of a small net income for the years ended December 31, 2002 and 1999, we have incurred net losses in each year since inception. Our losses have resulted principally from costs incurred in connection with our sales and marketing activities, research and development activities, manufacturing activities, general and administrative expenses associated with our operations, impairments on intangible assets and interest expense. Even if we are successful in launching additional products into the market, we expect to continue to incur substantial losses for the foreseeable future as we continue to sell and market our current products and research and develop, and seek regulatory approvals for, our product candidates.

If sales revenue from any of our current products or product candidates that receive marketing clearance from the FDA or other regulatory body is insufficient, if we are unable to develop and commercialize any of our product candidates, or if our product development is delayed, we may never become profitable. Even if we do become profitable, we may be unable to sustain or increase our profitability on a quarterly or annual basis.

Our success depends on our ability to successfully commercialize silicon nitride-based medical devices, which to date have experienced only limited market acceptance.

We believe we are the first and only company to use silicon nitride in medical applications. To date, however, we have had limited acceptance of our silicon nitride-based products and our product revenue has been derived substantially from our non-silicon nitride products. In order to succeed in our goal of becoming a leading biomaterial technology company utilizing silicon nitride, we must increase market awareness of our silicon nitride interbody spinal fusion products, continue to implement our sales and marketing strategy, enhance our commercial infrastructure and commercialize our silicon nitride joint replacement components and other products. If we fail in any of these endeavors or experience delays in pursuing them, we will not generate revenues as planned and will need to curtail operations or seek additional financing earlier than otherwise anticipated.

Our current products and our future products may not be accepted by hospitals and surgeons and may not become commercially successful.

Although we received 510(k) regulatory clearance from the FDA for our first silicon nitride spinal fusion products in 2008, we have not been able to obtain significant market share of the interbody spinal fusion market to date, and may not obtain such market share in the future. Even if we receive regulatory clearances or approvals for our product candidates in development, these product candidates may not gain market acceptance among orthopedic surgeons and the medical community. Orthopedic surgeons may elect not to use our products for a variety of reasons, including:

- lack or perceived lack of evidence supporting the beneficial characteristics of our silicon nitride technology;
- limited long-term data on the use of silicon nitride in medical devices;
- lower than expected clinical benefits in comparison with other products;
- the perception by surgeons that there are insufficient advantages of our products relative to currently available products;
- hospitals may choose not to purchase our products;
- group purchasing organizations may choose not to contract for our products, thus limiting availability of our products to hospital purchasers;
- the price of our products, which may be higher than products made of the other commonly used biomaterials in the interbody spinal fusion market and total joint market;

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- lack of coverage or adequate payment from managed care plans and other third-party payors for the procedures that use our products;
- Medicare, Medicaid or other third-party payors may limit or not permit reimbursement for procedures using our products;
- ineffective marketing and distribution support;
- the time and resources that may be required for training, or the inadequate training, of surgeons in the proper use of our products;
- the development of alternative biomaterials and products that render our products less competitive or obsolete; and
- the development of or improvement of competitive products.

If surgeons do not perceive our silicon nitride products and product candidates as superior alternatives to competing products, we will not be able to generate significant revenues, if any.

Even if surgeons are convinced of the superior characteristics of our silicon nitride products and our product candidates that we successfully introduce compared to the limitations of the current commonly used biomaterials, surgeons may find other methods or turn to other biomaterials besides silicon nitride to overcome such limitations. For instance, with respect to interbody spinal fusion products, surgeons or device manufacturers may use more effective markers for enhancing the imaging compatibility of PEEK devices, more effective antibiotics to prevent or treat implant-related infections, and more effective osteoconductive and osteoinductive materials when implanting an interbody spinal fusion device. Device manufacturers may also coat metal with existing traditional ceramics to reduce the risk of metal wear particles and corrosion in total joint replacement implants. Additionally, surgeons may increase their use of metal interbody spinal fusion devices if there is an increasing perception that PEEK devices are limited by their strength and resistance to fracture.

If we are unable to increase the productivity of our sales and marketing infrastructure we will not be able to penetrate the spinal fusion market.

We market and sell our products to surgeons and hospitals in the United States and select markets in Europe and South America using a network of independent third-party distributors who have existing surgeon relationships. We manage this distribution network through our in-house sales and marketing management team. We may also establish distribution collaborations in the United States and abroad in instances where access to a large or well-established sales and marketing organization may help to expand the market or accelerate penetration for selected products.

We cannot assure you that we will succeed in entering into and maintaining productive arrangements with an adequate number of distributors that are sufficiently committed to selling our products. The establishment of a distribution network is expensive and time consuming. As we launch new products and increase our marketing effort with respect to existing products, we will need to continue to hire, train, retain and motivate skilled independent distributors with significant technical knowledge in various areas, such as spinal fusion and total hip and knee joint replacement. In addition, the commissions we pay our distributors have increased over time, which has resulted in higher sales and marketing expenses, and those commissions and expenses may increase in the future. Furthermore, current and potential distributors may market and sell the products of our competitors. Even if the distributors market and sell our products, our competitors may be able, by offering higher commission payments or other incentives, to persuade these distributors to reduce or terminate their sales and marketing efforts related to our products. The distributors may also help competitors solicit business from our existing customers. Some of our independent distributors account for a significant portion of our sales volume, and, if we were to lose them, our sales could be adversely affected.

Even if we engage and maintain suitable relationships with an adequate number of distributors, they may not generate revenue as quickly as we expect them to, commit the necessary resources to effectively market and sell our products, or ultimately succeed in selling our products. We have been unable to obtain meaningful market share in the interbody spinal fusion device market with our current silicon nitride products to date and we may not be successful in increasing the productivity of our sales and marketing team and distribution network to gain meaningful market share for our silicon nitride products, which could adversely affect our business and financial condition.

The orthopedic market is highly competitive and we may not be able to compete effectively against the larger, well-established companies that dominate this market or emerging and small innovative companies that may seek to obtain or increase their share of the market.

The markets for spinal fusions and total hip and knee implant products are intensely competitive, and many of our competitors are much larger and have substantially more financial and human resources than we do. Many have long histories and strong reputations within the industry, and a relatively small number of companies dominate these markets. In 2012, Medtronic, Inc.; DePuy Synthes Companies, a group of Johnson & Johnson companies; Stryker Corporation; Biomet, Inc.; Zimmer Holdings, Inc.; and Smith & Nephew plc, accounted for more than 65% of orthopedic sales worldwide.

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These companies enjoy significant competitive advantages over us, including:

- broad product offerings, which address the needs of orthopedic surgeons and hospitals in a wide range of procedures;
- products that are supported by long-term clinical data;
- greater experience in, and resources for, launching, marketing, distributing and selling products, including strong sales forces and established distribution networks;
- existing relationships with spine and joint reconstruction surgeons;
- extensive intellectual property portfolios and greater resources for patent protection;
- greater financial and other resources for product research and development;
- greater experience in obtaining and maintaining FDA and other regulatory clearances and approvals for products and product enhancements;
- established manufacturing operations and contract manufacturing relationships;
- significantly greater name recognition and widely recognized trademarks; and
- established relationships with healthcare providers and payors.

Our products and any product candidates that we may introduce into the market may not enable us to overcome the competitive advantages of these large and dominant orthopedic companies. In addition, even if we successfully introduce additional product candidates incorporating our silicon nitride biomaterial into the market, emerging and small innovative companies may seek to increase their market share and they may eventually possess competitive advantages, which could adversely impact our business. Our competitors may also employ pricing strategies that could adversely affect the pricing of our products and pricing in the spinal fusion and total joint replacement market generally.

Moreover, many other companies are seeking to develop new biomaterials and products which may compete effectively against our products in terms of performance and price. For example, Smith & Nephew has developed a ceramic-coated metal, known as Oxinium, which may overcome certain of the limitations of metal joint replacement products and could directly compete with our silicon nitride and silicon nitride-coated product candidates.

We have significant customer concentration, so that economic difficulties or changes in the purchasing policies or patterns of our key customers could have a significant impact on our business and operating results.

A small number of customers account for a substantial portion of our product revenues. Our customers are primarily hospitals and surgical centers. At December 31, 2013, our largest customer, Bon Secours St. Mary's Hospital, or St. Mary's, had a receivable balance of approximately 15% of our total trade accounts receivable. In addition, St. Mary's accounted for 14% of our product revenues for each of the years ended December 31, 2013 and 2012. Sales of our products to our customers, including St. Mary's, are not based on long-term, committed-volume purchase contracts, and we may not continue to receive significant revenues from St. Mary's or any customer. Because of our significant customer concentration, our revenue could fluctuate significantly due to changes in economic conditions, the use of competitive products, or the loss of, reduction of business with, or less favorable terms with St. Mary's or any of our other significant customers. A significant portion of St. Mary's' purchases have been of our non-silicon nitride products, so it may be able to purchase competitive similar products from others. A reduction or delay in orders from St. Mary's or any of our other significant customers, or a delay or default in payment by any significant customer, could materially harm our business and results of operations.

The manufacturing process for our silicon nitride products is complex and requires sophisticated state-of-the-art equipment, experienced manufacturing personnel and highly specialized knowledge. If we are unable to manufacture our silicon nitride products on a timely basis consistent with our quality standards, our results of operation will be adversely impacted.

In order to control the quality, cost and availability of our silicon nitride products, we developed our own manufacturing capabilities. We operate a 30,000 square foot manufacturing facility which is certified under the ISO 13485 medical device manufacturing standard for medical devices and operates under the FDA's quality systems regulations, or QSRs. All operations with the exceptions of raw material production, cleaning, packaging and sterilization are performed at this facility.

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We currently do not have a secondary source for the manufacture of our silicon nitride products. Our reliance solely on our internal resources to manufacture our silicon nitride products entails risks to which we would not be subject if we had secondary suppliers for their manufacture, including:

- the inability to meet our product specifications and quality requirements consistently;
- a delay or inability to procure or expand sufficient manufacturing capacity to meet additional demand for our products;
- manufacturing and product quality issues related to the scale-up of manufacturing;
- the inability to produce a sufficient supply of our products to meet product demands;
- the disruption of our manufacturing facility due to equipment failure, natural disaster or failure to retain key personnel; and
- our inability to ensure our compliance with regulations and standards of the FDA including QSRs and corresponding state and international regulatory authorities.

Any of these events could lead to a reduction in our product sales, product launch delays, failure to obtain regulatory clearance or approval or impact our ability to successfully sell our products and commercialize our products candidates. While we currently are experiencing an equipment repair and have been able to obtain product from a supplier in the interim, we may be unable to do so in the future. Some of these events could be the basis for adverse actions by regulatory authorities, including injunctions, recalls, seizures, or total or partial suspension of production. In November 2013, we entered into a cooperative research and development agreement with Kyocera Industrial Ceramics Corporation, or Kyocera, under which we will work with Kyocera to determine its ability to become a second qualified manufacturer of our silicon nitride-based spinal fusion products and product candidates. Although we expect this arrangement will lead to Kyocera becoming a secondary qualified manufacturer, if Kyocera fails to become a qualified manufacturer or if we cannot come to an agreement with Kyocera for the further manufacture of our silicon nitride-based spinal fusion products and product candidates, we will continue to be the sole manufacturer of these products and will need to seek other potential secondary manufacturers.

We depend on a limited number of third-party suppliers for key raw materials used in the manufacturing of our silicon nitride products, and the loss of these third-party suppliers or their inability to supply us with adequate raw materials could harm our business.

We rely on a limited number of third-party suppliers for the raw materials required for the production of our silicon nitride products and product candidates. Our dependence on a limited number of third-party suppliers involves several risks, including limited control over pricing, availability, quality, and delivery schedules for raw materials. We have no supply agreements in place with any of our suppliers and cannot be certain that our current suppliers will continue to provide us with the quantities of raw materials that we require or that satisfy our anticipated specifications and quality requirements. Any supply interruption in limited or single sourced raw materials could materially harm our ability to manufacture our products until a new source of supply, if any, could be identified and qualified. We may be unable to find a sufficient alternative supply channel within a reasonable time or on commercially reasonable terms. Any performance failure on the part of our suppliers could delay the production of our silicon nitride products and product candidates and delay the development and commercialization of our product candidates, including limiting supplies necessary for commercial sale, clinical trials and regulatory approvals, which could have a material adverse effect on our business.

Use of third-party manufacturers increases the risk that we will not have adequate supplies of our non-silicon nitride products or instrumentation sets.

The majority of our product revenue is currently generated by sales of non-silicon nitride products. Our reliance on a limited number of third-party manufacturers to supply us with our non-silicon nitride products and instruments exposes us to risks that could delay our sales, or result in higher costs or lost product revenues. In particular, our manufacturers could:

- encounter difficulties in achieving volume production, quality control and quality assurance or suffer shortages of qualified personnel, which could result in their inability to manufacture sufficient quantities of our commercially available non-silicon nitride products to meet market demand for those products, or they could experience similar problems that result in the manufacture of insufficient quantities of our non-silicon nitride product candidates; and
- fail to follow and remain in compliance with the FDA-mandated QSRs, compliance which is required for all medical devices, or fail to document their compliance to QSRs, either of which could lead to significant delays in the availability of materials for our non-silicon nitride products or instrumentation sets.

If we are unable to obtain adequate supplies of our non-silicon nitride products and related instrumentation sets that meet our specifications and quality standards, it will be difficult for us to compete effectively. We have no supply agreements in place with our

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manufacturers and they may change the terms of our future orders or choose not to supply us with products or instrumentation sets in the future. Furthermore, if a third-party manufacturer from whom we purchase fails to perform its obligations, we may be forced to purchase products or related instrumentation from other third-party manufacturers, which we may not be able to do on reasonable terms, if at all. In addition, if we are required to change manufacturers for any reason, we will be required to verify that the new manufacturer maintains facilities and procedures that comply with quality standards and with all applicable regulations and guidelines. The delays associated with the verification of a new manufacturer or the re-verification of an existing manufacturer could negatively affect our ability to produce and distribute our non-silicon nitride products or instruments in a timely manner.

In order to be successful, we must expand our available product lines of silicon nitride-based medical devices by commercializing new product candidates, but we may not be able to do so in a timely fashion and at expected costs, or at all.

Although we are currently marketing our silicon nitride interbody spinal fusion implants, in order to be successful, we will need to expand our product lines to include other silicon nitride devices. Therefore, we are developing silicon nitride product candidates for total hip and knee replacement procedures and are exploring the application of our silicon nitride technology for other potential applications. However, we have yet to commercialize any silicon nitride products beyond our spinal fusion products. To succeed in our commercialization efforts, we must effectively continue product development and testing, obtain regulatory clearances and approvals, and enhance our sales and marketing capabilities. We may also have to write down significant inventory if existing products are replaced by new products. Because of these uncertainties, there is no assurance that we will succeed in bringing any of our current or future product candidates to market. If we fail in bringing our product candidates to market, or experience delays in doing so, we will not generate revenues as planned and will need to curtail operations or seek additional financing earlier than otherwise anticipated.

We will depend on one or more strategic partners to develop and commercialize our total joint replacement product candidates, and if our strategic partners are unable to execute effectively on our agreements with them, we may never become profitable.

Pursuant to a joint development and license agreement with Orthopaedic Synergy, Inc., or OSI, we are dependent on OSI's ability to execute product development plans, obtain regulatory approvals, and sell, distribute and market our jointly developed product candidate for total hip and total knee joint replacement implants that use our *MC²* silicon nitride technology. We would similarly be reliant on other strategic partners to develop and commercialize a total hip or knee joint replacement product candidate that utilizes silicon nitride-coated components, although we have not yet entered into an agreement with any strategic partner to develop products with these silicon nitride-coated components and may be unable to do so on agreeable terms. In order to succeed in our joint commercialization efforts, we and OSI, and any future partners must execute effectively on all elements of a combined business plan, including continuing to establish sales and marketing capabilities, manage certified, validated and effective commercial-scale manufacturing operations, conduct product development and testing, and obtain regulatory clearances and approvals for our product candidate. If we or any of our strategic partners fail in any of these endeavors, or experience delays in pursuing them, we will not generate revenues as planned and will need to curtail operations or seek additional financing earlier than otherwise anticipated.

The use of physician-owned distributorships could result in increased pricing pressure on our products or harm our ability to sell our products to physicians who own or are affiliated with those distributorships and the sale of our products through such distributorships may expose us to regulatory enforcement risk.

Physician-owned distributorships, or PODs, are medical device distributors that are owned, directly or indirectly, by physicians. These physicians derive a proportion of their revenue from selling or arranging for the sale of medical devices for use in procedures they perform on their own patients at hospitals that agree to purchase from or through the POD, or that otherwise furnish ordering physicians with income that is based directly or indirectly on those orders of medical devices.

We may sell and distribute our products through a limited number of PODs. The number of PODs in the orthopedic industry may continue to grow as physicians search for ways to increase their incomes. These companies and the physicians who own, or partially own, them have significant market knowledge and access to the surgeons and hospitals that may potentially purchase our products and the physicians who own these PODs will have financial incentives to purchase from these distributorships. As a result, growth in this area may reduce our ability to compete effectively for business.

On March 26, 2013, the Department of Health and Human Services Office of Inspector General issued a Special Fraud Alert on Physician-Owned Entities and identified PODs as "inherently suspect" under the federal Anti-Kickback Statute. While the PODs themselves may be the target of any government enforcement efforts in this area, it is possible that regulatory scrutiny may extend to other entities that have relationships with PODs, including us. We are not aware that we are currently subject to any such scrutiny. However, the cost of defending such enforcement actions, if brought (even without merit), as well as any sanctions, if imposed, could have a material adverse effect on our business.

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If hospitals and other healthcare providers are unable to obtain coverage or adequate reimbursement for procedures performed with our products, it is unlikely our products will be widely used.

In the United States, the commercial success of our existing products and any future products will depend, in part, on the extent to which governmental payors at the federal and state levels, including Medicare and Medicaid, private health insurers and other third-party payors provide coverage for and establish adequate reimbursement levels for procedures utilizing our products. Because we typically receive payment directly from hospitals and surgical centers, we do not anticipate relying directly on payment from third-party payors for our products. However, hospitals and other healthcare providers that purchase our orthopedic products for treatment of their patients generally rely on third-party payors to pay for all or part of the costs and fees associated with our products as part of a “bundled” rate for the associated procedures. The existence of coverage and adequate reimbursement for our products and the procedures performed with them by government and private payors is critical to market acceptance of our existing and future products. Neither hospitals nor surgeons are likely to use our products if they do not receive adequate reimbursement for the procedures utilizing our products.

Many private payors currently base their reimbursement policies on the coverage decisions and payment amounts determined by the Centers for Medicare and Medicaid Services, or CMS, which administers the Medicare program. Others may adopt different coverage or reimbursement policies for procedures performed with our products, while some governmental programs, such as Medicaid, have reimbursement policies that vary from state to state, some of which may not pay for the procedures performed with our products in an adequate amount, if at all. A Medicare national or local coverage decision denying coverage for one or more of our products could result in private and other third-party payors also denying coverage for our products. Third-party payors also may deny reimbursement for our products if they determine that a product used in a procedure was not medically necessary, was not used in accordance with cost-effective treatment methods, as determined by the third-party payor, or was used for an unapproved use. Unfavorable coverage or reimbursement decisions by government programs or private payors underscore the uncertainty that our products face in the market and could have a material adverse effect on our business.

Many hospitals and clinics in the United States belong to group purchasing organizations, which typically incentivize their hospital members to make a relatively large proportion of purchases from a limited number of vendors of similar products that have contracted to offer discounted prices. Such contracts often include exceptions for purchasing certain innovative new technologies, however. Accordingly, the commercial success of our products may also depend to some extent on our ability to either negotiate favorable purchase contracts with key group purchasing organizations and/or persuade hospitals and clinics to purchase our product “off contract.”

The healthcare industry in the United States has experienced a trend toward cost containment as government and private payors seek to control healthcare costs by paying service providers lower rates. While it is expected that hospitals will be able to obtain coverage for procedures using our products, the level of payment available to them for such procedures may change over time. State and federal healthcare programs, such as Medicare and Medicaid, closely regulate provider payment levels and have sought to contain, and sometimes reduce, payment levels. Private payors frequently follow government payment policies and are likewise interested in controlling increases in the cost of medical care. In addition, some payors are adopting pay-for-performance programs that differentiate payments to healthcare providers based on the achievement of documented quality-of-care metrics, cost efficiencies, or patient outcomes. These programs are intended to provide incentives to providers to deliver the same or better results while consuming fewer resources. As a result of these programs, and related payor efforts to reduce payment levels, hospitals and other providers are seeking ways to reduce their costs, including the amounts they pay to medical device manufacturers. We may not be able to sell our implants profitably if third-party payors deny or discontinue coverage or reduce their levels of payment below that which we project, or if our production costs increase at a greater rate than payment levels. Adverse changes in payment rates by payors to hospitals could adversely impact our ability to market and sell our products and negatively affect our financial performance.

In international markets, medical device regulatory requirements and healthcare payment systems vary significantly from country to country, and many countries have instituted price ceilings on specific product lines. We cannot assure you that our products will be considered cost-effective by international third-party payors, that reimbursement will be available or, if available, that the third-party payors’ reimbursement policies will not adversely affect our ability to sell our products profitably. Any failure to receive regulatory or reimbursement approvals would negatively impact market acceptance of our products in any international markets in which those approvals are sought.

Prolonged negative economic conditions in domestic and international markets may adversely affect us, our suppliers, partners and consumers, and the global orthopedic market which could harm our financial position.

Global credit and financial markets have been experiencing extreme disruptions over the past several years, including severely diminished liquidity and availability of credit, declines in consumer confidence, declines in economic growth, increases in unemployment rates and uncertainty about economic stability. Credit and financial markets and confidence in economic conditions

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might deteriorate further. Our business may be adversely affected by the recent economic downturn and volatile business environment and continued unpredictable and unstable market conditions. In addition, there is a risk that one or more of our current suppliers may not continue to operate. Any lender that is obligated to provide funding to us under any future credit agreement with us may not be able to provide funding in a timely manner, or at all, when we require it. The cost of, or lack of, available credit or equity financing could impact our ability to develop sufficient liquidity to maintain or grow our company. These negative changes in domestic and international economic conditions or additional disruptions of either or both of the financial and credit markets may also affect third-party payors and may have a material adverse effect on our business, results of operations, financial condition and liquidity.

In addition, we believe that various demographics and industry-specific trends will help drive growth in the orthopedics markets, but these demographics and trends are uncertain. Actual demand for orthopedic products generally, and our products in particular, could be significantly less than expected if our assumptions regarding these factors prove to be incorrect or do not materialize, or if alternative treatments gain widespread acceptance.

We have a new senior management team and are dependent on our senior management team, engineering team, sales and marketing team and surgeon advisors, and the loss of any of them could harm our business.

We have recently assembled a new senior management team. They have worked together in their new positions with us for a limited time and may not be able to successfully implement our strategy. In addition, we have not entered into employment agreements, other than severance agreements, with any of the members of our senior management team. There are no assurances that the services of any of these individuals will be available to us for any specified period of time. The successful integration of our new senior management team, the loss of members of our senior management team, sales and marketing team, engineering team and key surgeon advisors, or our inability to attract or retain other qualified personnel or advisors could have a material adverse effect on our business, financial condition and results of operations.

If we experience significant disruptions in our information technology systems, our business, results of operations and financial condition could be adversely affected.

The efficient operation of our business depends on our information technology systems. We rely on our information technology systems to effectively manage our sales and marketing, accounting and financial functions; manufacturing processes; inventory; engineering and product development functions; and our research and development functions. As such, our information technology systems are vulnerable to damage or interruption including from earthquakes, fires, floods and other natural disasters; terrorist attacks and attacks by computer viruses or hackers; power losses; and computer systems, or Internet, telecommunications or data network failures. The failure of our information technology systems to perform as we anticipate or our failure to effectively implement new systems could disrupt our entire operation and could result in decreased sales, increased overhead costs, excess inventory and product shortages, all of which could have a material adverse effect on our reputation, business, results of operations and financial condition.

Risks Related to Our Capital Resources and Impairments

We will require additional financing and our failure to obtain additional funding could force us to delay, reduce or eliminate our product development programs or commercialization efforts.

We require substantial future capital in order to continue to conduct the research and development and regulatory clearance and approval activities necessary to bring our products to market, to establish effective marketing and sales capabilities and to satisfy our obligations under our senior secured credit facility. Our existing capital resources, including the net proceeds from our initial public offering of shares of our common stock are not sufficient to enable us to fund the completion of the development and commercialization of all of our product candidates. We cannot determine with certainty the duration and completion costs of the current or future development and commercialization of our product candidates for spinal fusion procedures, joint replacement and coated metals or if, when, or to what extent we will generate revenues from the commercialization and sale of any of these product candidates for which we obtain regulatory approval. We may never succeed in achieving regulatory approval for certain of these product candidates. The duration, costs and timing of clinical trials and development of our spinal fusion, joint replacement and coated metal product candidates will depend on a variety of factors, including:

- the scope, rate of progress, and expense of our ongoing, as well as any additional, clinical trials and other research and development activities;
- future clinical trial results we may have to conduct;
- potential changes in government regulation; and
- the timing and receipt of any regulatory approvals.

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A change in the outcome of any of these variables with respect to the development of spinal fusion, joint replacement or coated metal product candidates could mean a significant change in the costs and timing associated with the development of these product candidates.

We currently have limited committed sources of capital and we have limited liquidity. Our cash and cash equivalents as of December 31, 2013 were \$2.3 million and as of December 31, 2012 were \$2.7 million. In December 2012, we entered into a senior secured credit facility with General Electric Capital Corporation, or GE Capital, as agent and lender, and Zions First National Bank, as lender, which is described in more detail in the “Management’s Discussion and Analysis of Financial Condition and Results of Operations” section of this report and which we refer to as the GE Secured Lending Facility. The GE Secured Lending Facility consists of an \$18.0 million 30-month term loan and a \$3.5 million revolving credit facility. The revolving line of credit is secured by our accounts receivable, based on certain defined criteria. In January 2014, we began making monthly repayment of the principal amount due under the term loan of \$600,000 per month. Due to the amortization of our term loan, we expect to use a substantial amount of our monthly cash flow to repay the GE Secured Lending Facility.

The GE Secured Lending Facility contains certain financial covenants related to monthly cash burn, as defined in the revolving credit facility, minimum liquidity, days sales outstanding of accounts receivable balances, annual payment restrictions to certain company affiliates and other financial reporting requirements. Specifically, under the liquidity covenant in the revolving credit facility, we are required to maintain cash and cash equivalents and availability under the GE Secured Lending Facility of equal to the greater of \$1.5 million (exclusive of availability under the revolving credit facility) or six times our monthly cash burn. We were in default of this liquidity covenant in November 2013. In December 2013, we obtained a waiver of this liquidity covenant from November 1, 2013 through January 31, 2014 and agreed to increase the credit reserve under this facility from \$0.5 million to \$1.0 million. On January 28, 2014, we obtained an additional waiver of the liquidity covenant from GE Capital through February 28, 2014 and agreed to increase the credit reserve under this facility by an additional \$0.5 million, bringing the total reserve to \$1.5 million. We were in compliance with the liquidity covenant on February 28, 2014. However, we will need to obtain additional funding during the third quarter of 2014 to satisfy the \$7.2 million of principal due under the GE Secured Lending Facility in 2014 (which amount excludes interest payments and any other fees and charges incurred or that we may incur) and to maintain compliance with the financial and liquidity covenants related to the GE Secured Lending Facility through 2014. Furthermore, if we are unable to access additional funds prior to becoming non-compliant on any of the financial and liquidity covenants, the entire remaining balance of the GE Secured Lending Facility could become immediately due and payable at the option of GE Capital.

In addition, the repayment of the GE Secured Lending Facility and the liquidity covenant limit our ability to use our cash and cash equivalents to fund our operations and may restrict our ability to continue development of our product candidates. Additionally, the GE Secured Lending Facility restricts our ability to incur additional *pari passu* indebtedness, which may reduce our ability to seek additional financing. If adequate funds are not available on a timely basis, we may terminate or delay the development of one or more of our product candidates, or delay activities necessary to commercialize our product candidates. Additional funding may not be available to us on acceptable terms, or at all. Any additional equity financing, if available, may not be available on favorable terms and will most likely be dilutive to our current stockholders, and debt financing, if available, may involve more restrictive covenants. Our ability to access capital when needed is not assured and, if not achieved on a timely basis, will materially harm our business, financial condition and results of operations.

As a result of our debt obligations, we will need additional funds to meet our operational needs and capital requirements for product development, clinical trials and commercialization. The timing and amount of our future capital requirements will depend on many factors, including:

- our ability to satisfy our obligation to pay principal, interest and fees on the GE Secured Lending Facility;
- our ability to comply with covenants related to the GE Secured Lending Facility;
- the level of sales of our current products and the cost of revenue and sales and marketing;
- the extent of any clinical trials that we will be required to conduct in support of the regulatory clearance of our total hip and knee replacement product candidates;
- the scope, progress, results and cost of our product development efforts;
- the costs, timing and outcomes of regulatory reviews of our product candidates;
- the number and types of products we develop and commercialize;
- the costs of preparing, filing and prosecuting patent applications and maintaining, enforcing and defending intellectual property-related claims; and
- the extent and scope of our general and administrative expenses.

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Raising additional capital by issuing securities or through debt financings or licensing arrangements may cause dilution to existing stockholders, restrict our operations or require us to relinquish proprietary rights.

To the extent that we raise additional capital through the sale of equity or convertible debt securities, your ownership interest will be diluted, and the terms may include liquidation or other preferences that adversely affect your rights as a stockholder. Debt financing, if available, may involve agreements that include covenants limiting or restricting our ability to take specific actions such as incurring additional debt, making capital expenditures or declaring dividends. If we raise additional funds through collaboration and licensing arrangements with third parties, we may have to relinquish valuable rights to our technologies or products or grant licenses on terms that are not favorable to us. Any of these events could adversely affect our ability to achieve our product development and commercialization goals and have a material adverse effect on our business, financial condition and results of operations.

Our independent registered public accounting firm has included an explanatory paragraph relating to our ability to continue as a going concern in its report on our audited financial statements. We may be unable to continue to operate without the threat of liquidation for the foreseeable future.

Our report from our independent registered public accounting firm for the year ended December 31, 2013 includes an explanatory paragraph stating that our recurring losses from operations and our need to obtain additional financing in order to satisfy our debt obligations and to be compliant with covenants under our debt obligations through 2014 raise substantial doubt about our ability to continue as a going concern. If we are unable to obtain sufficient funding, our business, prospects, financial condition and results of operations will be materially and adversely affected and we may be unable to continue as a going concern. If we are unable to continue as a going concern, we may have to liquidate our assets and may receive less than the value at which those assets are carried on our consolidated financial statements, and it is likely that investors will lose all or a part of their investment. Future reports from our independent registered public accounting firm may also contain statements expressing doubt about our ability to continue as a going concern. If we seek additional financing to fund our business activities in the future and there remains doubt about our ability to continue as a going concern, investors or other financing sources may be unwilling to provide additional funding on commercially reasonable terms or at all.

If we do not adhere to the financial covenants set forth in our GE Secured Lending Facility, we will be in default of our GE Secured Lending Facility.

The GE Secured Lending Facility includes certain financial covenants including a requirement that the average time that it takes us to collect on any amounts due to us from any customers not exceed 85 days for any calendar month, as well as a liquidity covenant. The liquidity covenant may significantly limit our ability to use our cash as cash equivalents to fund our operations as it requires us to maintain cash and cash equivalents and availability under the revolving credit facility equal to the greater of \$1.5 million (exclusive of availability under the revolving credit facility) or six times our monthly cash burn, as defined in the facility. As of December 31, 2013, six times our monthly cash burn equaled \$4.0 million. We were in default of this liquidity covenant in November 2013. In December 2013, we obtained a waiver of this liquidity covenant from November 1, 2013 through January 31, 2014 and agreed to increase the credit reserve under this facility from \$0.5 million to \$1.0 million. On January 28, 2014, we obtained an additional waiver of the liquidity covenant from GE Capital through February 28, 2014 and agreed to increase the credit reserve under this facility by an additional \$0.5 million, bringing the total reserve to \$1.5 million.

We are seeking to refinance the GE Secured Lending Facility or obtain additional financing. However, we may have difficulty obtaining additional debt financing, due to the restrictions in the GE Secured Lending Facility and may have difficulty in refinancing the facility. There is no guarantee we will be successful in entering into any such lending arrangement on commercially reasonable terms, or at all. Moreover, even if we are able to enter into a new lending arrangement sufficient to repay the GE Secured Lending Facility, such new facility will likely contain liquidity, financial and operational covenants, which could be as restrictive as or more restrictive than those in the GE Secured Lending Facility. In addition, even if we are successful in obtaining additional financing, the terms of such additional debt could further restrict our operating and financial flexibility. Further, if we are liquidated, the lenders' right to repayment would be senior to the rights of the holders of our common stock to receive any proceeds from the liquidation. The agent could declare a default under the GE Secured Lending Facility upon the occurrence of a material adverse effect, as defined under the loan agreement, thereby requiring us to either repay the outstanding indebtedness immediately or attempt to reverse the declaration of default through negotiation or litigation. Any declaration by the agent of an event of default could significantly harm our business and prospects and could cause the price of our common shares to decline.

An impairment charge could have a material adverse effect on our financial condition and results of operations.

We are required to test acquired goodwill for impairment on an annual basis. Goodwill represents the excess of the amount paid over the fair value of the net assets at the date of the acquisition. We have chosen to complete our annual impairment reviews of goodwill at the end of each calendar year. We also are required to test goodwill for impairment between annual tests if events occur or

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circumstances change that would more likely than not reduce our enterprise fair value below its book value. In addition, we are required to test our finite-lived intangible assets for impairment if events occur or circumstances change that would indicate the remaining net book value of the finite-lived intangible assets might not be recoverable. These events or circumstances could include a significant change in the business climate, including a significant sustained decline in our market value, legal factors, operating performance indicators, competition, sale or disposition of a significant portion of our business and other factors.

If the fair market value of our reporting unit is less than its book value, we could be required to record an impairment charge. The valuation of a reporting unit requires judgment in estimating future cash flows, discount rates and other factors. In making these judgments, we evaluate the financial health of our business, including such factors as industry performance, changes in technology and operating cash flows. Changes in our forecasts or decreases in the value of our common stock could cause book values of our reporting unit to exceed its fair value, which may result in goodwill impairment charges. The amount of any impairment could be significant and could have a material adverse effect on our reported financial results for the period in which the charge is taken.

Risks Related to Regulatory Approval of Our Products and Other Government Regulations

Our long-term success depends substantially on our ability to obtain regulatory clearance or approval and thereafter commercialize our product candidates; we cannot be certain that we will be able to do so in a timely manner or at all.

The process of obtaining regulatory clearances or approvals to market a medical device from the FDA or similar regulatory authorities outside of the United States can be costly and time consuming, and there can be no assurance that such clearances or approvals will be granted on a timely basis, or at all. The FDA's 510(k) clearance process generally takes one to six months from the date of submission, depending on whether a special or traditional 510(k) premarket notification has been submitted, but can take significantly longer. An application for premarket approval, or PMA, must be submitted to the FDA if the device cannot be cleared through the 510(k) clearance process or is not exempt from premarket review by the FDA. The PMA process almost always requires one or more clinical trials and can take two to three years from the date of filing, or even longer. In some cases, including in the case of our interbody spinal fusion devices which incorporate our *CSC* technology and our *MC²* silicon nitride femoral head component, the FDA requires clinical data as part of the 510(k) clearance process.

It is possible that the FDA could raise questions about our spinal fusion products, our spinal fusion product candidates and our total hip and knee joint replacement product candidates and could require us to perform additional studies on our products and product candidates. Even if the FDA permits us to use the 510(k) clearance process, we cannot assure you that the FDA will not require either supporting data from laboratory tests or studies that we have not conducted, or substantial supporting clinical data. If we are unable to use the 510(k) clearance process for any of our product candidates, are required to provide clinical data or laboratory data that we do not possess to support our 510(k) premarket notifications for any of these product candidates, or otherwise experience delays in obtaining or fail to obtain regulatory clearances, the commercialization of our product candidates in the United States will be delayed or prevented, which will adversely affect our ability to generate additional revenues. It also may result in the loss of potential competitive advantages that we might otherwise attain by bringing our products to market earlier than our competitors. Additionally, although the FDA allows modifications to be made to devices that have received 510(k) clearance with supporting documentation, the FDA may disagree with our decision to modify our cleared devices without submission of a new 510(k) premarket notification, subjecting us to potential product recall, field alerts and corrective actions. Any of these contingencies could adversely affect our business.

Similar to our compliance with U.S. regulatory requirements, we must obtain and comply with international requirements in order to market and sell our products outside of the United States and we may only promote and market our products, if approved, as permitted by applicable regulatory authorities.

The safety of our products is not yet supported by long-term clinical data, and they may prove to be less safe and effective than our laboratory data indicate.

We obtained FDA clearance for each of our products that we currently market, and we have sought and intend to seek FDA clearance or approval through the FDA's 510(k) or PMA process and, where applicable, CE marking for our product candidates. The 510(k) clearance process is based on the FDA's agreement that a new product candidate is substantially equivalent to an already marketed product for which a PMA was not required. While most 510(k) premarket notifications do not require clinical data for clearance, the FDA may request that such data be provided. Long-term clinical data or marketing experience obtained after clearance may indicate that our products cause unexpected complications or other unforeseen negative effects. If this happens, we could be subject to the withdrawal of our marketing clearance and other enforcement sanctions by the FDA or other regulatory authority, product recalls, significant legal liability, significant negative publicity, damage to our reputation and a dramatic reduction in our ability to sell our products, any one of which would have a material adverse effect on our business, financial condition and results of operations.

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We expect to be required to conduct clinical trials to support regulatory approval of some of our product candidates. We have no experience conducting clinical trials, they may proceed more slowly than anticipated, and we cannot be certain that our product candidates will be shown to be safe and effective for human use.

In order to commercialize our product candidates in the United States, we must submit a PMA for some of these product candidates, which will require us to conduct clinical trials. We also plan to provide the FDA with clinical trial data to support some of our 510(k) premarket notifications. We will receive approval or clearance from the FDA to commercialize products requiring a clinical trial only if we can demonstrate to the satisfaction of the FDA, through well-designed and properly conducted clinical trials, that our product candidates are safe and effective and otherwise meet the appropriate standards required for approval or clearance for specified indications. Clinical trials are complex, expensive, time consuming, uncertain and subject to substantial and unanticipated delays. Before we may begin clinical trials, we must submit and obtain approval for an investigational device exemption, or IDE, that describes, among other things, the manufacture of, and controls for, the device and a complete investigational plan. Clinical trials generally involve a substantial number of patients in a multi-year study. Because we do not have the experience or the infrastructure necessary to conduct clinical trials, we will have to hire one or more contract research organizations, or CROs, to conduct trials on our behalf. CRO contract negotiations may be costly and time consuming and we will rely heavily on the CRO to ensure that our trials are conducted in accordance with regulatory and industry standards. We may encounter problems with our clinical trials and any of those problems could cause us or the FDA to suspend those trials, or delay the analysis of the data derived from them.

A number of events or factors, including any of the following, could delay the completion of our clinical trials in the future and negatively impact our ability to obtain FDA approval for, and to introduce our product candidates:

- failure to obtain financing necessary to bear the cost of designing and conducting clinical trials;
- failure to obtain approval from the FDA or foreign regulatory authorities to commence investigational studies;
- conditions imposed on us by the FDA or foreign regulatory authorities regarding the scope or design of our clinical trials;
- failure to find a qualified CRO to conduct our clinical trials or to negotiate a CRO services agreement on favorable terms;
- delays in obtaining or in our maintaining required approvals from institutional review boards or other reviewing entities at clinical sites selected for participation in our clinical trials;
- insufficient supply of our product candidates or other materials necessary to conduct our clinical trials;
- difficulties in enrolling patients in our clinical trials;
- negative or inconclusive results from clinical trials, or results that are inconsistent with earlier results, that necessitate additional clinical studies;
- failure on the part of the CRO to conduct the clinical trial in accordance with regulatory requirements;
- our failure to maintain a successful relationship with the CRO or termination of our contractual relationship with the CRO before completion of the clinical trials;
- serious or unexpected side effects experienced by patients in whom our product candidates are implanted; or
- failure by any of our third-party contractors or investigators to comply with regulatory requirements or meet other contractual obligations in a timely manner.

Our clinical trials may need to be redesigned or may not be completed on schedule, if at all. Delays in our clinical trials may result in increased development costs for our product candidates, which could cause our stock price to decline and limit our ability to obtain additional financing. In addition, if one or more of our clinical trials are delayed, competitors may be able to bring products to market before we do, and the commercial viability of our product candidates could be significantly reduced.

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Our current and future relationships with third-party payors and current and potential customers in the United States and elsewhere may be subject, directly or indirectly, to applicable anti-kickback, fraud and abuse, false claims, transparency, health information privacy and security and other healthcare laws and regulations, which could expose us to criminal sanctions, civil penalties, contractual damages, reputational harm administrative burdens and diminished profits and future earnings.

Our current and future arrangements with third-party payors and current and potential customers, including providers and physicians, as well as PODs, as discussed above, may expose us to broadly applicable fraud and abuse and other healthcare laws and regulations, including, without limitation, the federal Anti-Kickback Statute and the federal False Claims Act, which may constrain the business or financial arrangements and relationships through which we sell, market and distribute our products. In addition, we may be subject to transparency laws and patient privacy regulations by U.S. federal and state governments and by governments in foreign jurisdictions in which we conduct our business. The applicable federal, state and foreign healthcare laws and regulations that may affect our ability to operate include:

- the federal Anti-Kickback Statute, which prohibits, among other things, persons from knowingly and willfully soliciting, offering, receiving or providing remuneration, directly or indirectly, in cash or in kind, to induce or reward, or in return for, either the referral of an individual for, or the purchase, order or recommendation of, any good or service, for which payment may be made under federal healthcare programs, such as Medicare and Medicaid;
- federal civil and criminal false claims laws and civil monetary penalty laws, including the federal False Claims Act, which impose criminal and civil penalties, including civil whistleblower or qui tam actions, against individuals or entities for knowingly presenting, or causing to be presented, to the federal government, including the Medicare and Medicaid programs, claims for payment that are false or fraudulent or making a false statement to avoid, decrease or conceal an obligation to pay money to the federal government;
- the federal Health Insurance Portability and Accountability Act of 1996, or HIPAA, which imposes criminal and civil liability for executing a scheme to defraud any healthcare benefit program or making false statements relating to healthcare matters;
- HIPAA, as amended by the Health Information Technology for Economic and Clinical Health Act of 2009, or HITECH, and their respective implementing regulations, which impose obligations on covered healthcare providers, health plans, and healthcare clearinghouses, as well as their business associates that create, receive, maintain or transmit individually identifiable health information for or on behalf of a covered entity, with respect to safeguarding the privacy, security and transmission of individually identifiable health information;
- the Physician Payments Sunshine Act, which requires (i) manufacturers of drugs, devices, biologics and medical supplies for which payment is available under Medicare, Medicaid or the Children's Health Insurance Program, with specific exceptions, to report annually to CMS information related to certain "payments or other transfers of value" made to physicians, which is defined to include doctors, dentists, optometrists, podiatrists and chiropractors, and teaching hospitals, with data collection beginning on August 1, 2013, (ii) applicable manufacturers and applicable group purchasing organizations to report annually to CMS ownership and investment interests held in such entities by physicians and their immediate family members, with data collection beginning on August 1, 2013, (iii) manufacturers to submit reports to CMS by March 31, 2014 and the 90th day of each subsequent calendar year, and (iv) disclosure of such information by CMS on a publicly available website beginning in September 2014; and
- analogous state and foreign laws and regulations, such as state anti-kickback and false claims laws, which may apply to sales or marketing arrangements and claims involving healthcare items or services reimbursed by non-governmental third-party payors, including private insurers; state and foreign laws that require medical device companies to comply with the medical device industry's voluntary compliance guidelines and the relevant compliance guidance promulgated by the federal government or otherwise restrict payments that may be made to healthcare providers; state and foreign laws that require medical device manufacturers to report information related to payments and other transfers of value to physicians and other healthcare providers or marketing expenditures; and state and foreign laws governing the privacy and security of health information in certain circumstances, many of which differ from each other in significant ways and often are not preempted by HIPAA, thus complicating compliance efforts.

Efforts to ensure that our business arrangements with third parties will comply with applicable healthcare laws and regulations may involve substantial costs. It is possible that governmental authorities will conclude that our business practices may not comply with current or future statutes, regulations or case law involving applicable fraud and abuse or other healthcare laws and regulations. If our operations are found to be in violation of any of these laws or any other governmental regulations that may apply to us, we may be subject to significant civil, criminal and administrative penalties, including, without limitation, damages, fines, imprisonment, exclusion from participation in government healthcare programs, such as Medicare and Medicaid, and the curtailment or restructuring of our operations, which could have a material adverse effect on our business. If any of the physicians or other healthcare providers or entities with whom we expect to do business, including our collaborators, are found not to be in compliance with applicable laws, they may be subject to criminal, civil or administrative sanctions, including exclusions from participation in government healthcare programs, which could also materially affect our business.

In July 2012, we received a subpoena from the Department of Justice seeking the production of documents, including documents related to our relationship with a particular customer and various entities, including a company distributor, and individuals associated with that distributor. In April 2013, we received a second subpoena requesting similar records. We cooperated with the Department of Justice's requests and provided the records requested by the two subpoenas. We have had no further communications with the Department of Justice since responding to its second request in June 2013. While we do not believe that we are the target of the government's investigation, if we are found to have violated one or more applicable laws, we could be subject to the risks and consequences discussed above. In addition, responding to any additional requests or actions of the Department of Justice in connection with this investigation may be expensive and time-consuming.

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Recently enacted and future legislation may increase the difficulty and cost for us to obtain regulatory approval or clearance of our product candidates and affect the prices we may obtain for our products.

In the United States and some foreign jurisdictions, there have been a number of legislative and regulatory changes and proposed changes regarding the healthcare system that could prevent or delay clearance and/or approval of our product candidates, restrict or regulate post-clearance and post-approval activities and affect our ability to profitably sell our products and any product candidates for which we obtain marketing approval or clearance.

In addition, FDA regulations and guidance are often revised or reinterpreted by the FDA in ways that may significantly affect our business and our products. Any new regulations or revisions or reinterpretations of existing regulations may impose additional costs or lengthen review times of our products. Delays in receipt of or failure to receive regulatory clearances or approvals for our new products would have a material adverse effect on our business, results of operations and financial condition. In addition, the FDA is currently evaluating the 510(k) process and may make substantial changes to industry requirements, including which devices are eligible for 510(k) clearance, the ability to rescind previously granted 510(k) clearances and additional requirements that may significantly impact the process.

Among policy makers and payors in the United States and elsewhere, there is significant interest in promoting changes in healthcare systems with the stated goals of containing healthcare costs, improving quality and expanding access. In the United States, the medical device industry has been a particular focus of these efforts and has been significantly affected by major legislative initiatives. In March 2010, President Obama signed into law the Patient Protection and Affordable Care Act, as amended by the Health Care and Education Affordability Reconciliation Act, or collectively the ACA, a sweeping law intended, among other things, to broaden access to health insurance, reduce or constrain the growth of healthcare spending, enhance remedies against fraud and abuse, add new transparency requirements for the healthcare and health insurance industries, impose new taxes and fees on the health industry and impose additional health policy reforms.

Among the provisions of the ACA of importance to our products and product candidates are:

- a 2.3% medical device excise tax on the U.S. sales of most medical devices, which currently includes and we expect will continue to include U.S. sales of our current products and product candidates that receive clearance or approval;
- expansion of healthcare fraud and abuse laws, including the False Claims Act and the Anti-Kickback Statute, and new government investigative powers and enhanced penalties for non-compliance;
- new requirements under the federal Open Payments program and its implementing regulations;
- a new Patient-Centered Outcomes Research Institute to oversee, identify priorities in, and conduct comparative clinical effectiveness research, along with funding for such research; and
- creation of an independent payment advisory board that will submit recommendations to reduce Medicare spending if projected Medicare spending exceeds a specified growth rate.

In addition, other legislative changes have been proposed and adopted since the ACA was enacted. For example, on August 2, 2011, the President signed into law the Budget Control Act of 2011, which, among other things, created the Joint Select Committee on Deficit Reduction to recommend to Congress proposals in spending reductions. The Joint Select Committee on Deficit Reduction did not achieve a targeted deficit reduction of at least \$1.2 trillion for the years 2013 through 2021, triggering the legislation's automatic reduction to several government programs. This includes aggregate reductions to Medicare payments to providers of up to 2% per fiscal year, starting in 2013. On January 2, 2013, President Obama signed into law the American Taxpayer Relief Act of 2012, or ATRA, which, among other things, reduced Medicare payments to several types of providers and increased the statute of limitations period for the government to recover overpayments to providers from three to five years. On March 1, 2013, the President signed an executive order implementing the Budget Control Act's 2% Medicare payment reductions, and on April 1, 2013, these reductions went into effect. These new laws may result in additional reductions in Medicare and other healthcare funding, which could have a material adverse effect on our financial operations.

We expect that the ACA, as well as other healthcare reform measures that have been and may be adopted in the future, may result in more rigorous coverage criteria and in additional downward pressure on the price that we receive for our products. Any reduction in reimbursement from Medicare or other government programs may result in a similar reduction in payments from private payors. The implementation of cost containment measures or other healthcare reforms may affect our ability to generate revenue and profits or commercialize our product candidates.

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In the European Union and some other international markets, the government provides health care at a low cost to consumers and regulates prices of healthcare products, patient eligibility or reimbursement levels to control costs for the government-sponsored health care system. Many countries are reducing their public expenditures and we expect to see strong efforts to reduce healthcare costs in international markets, including patient access restrictions, suspensions on price increases, prospective and possibly retroactive price reductions and other recoupments and increased mandatory discounts or rebates and recoveries of past price increases. These cost control measures could reduce our revenues. In addition, certain countries set prices by reference to the prices in other countries where our products are marketed. Thus, our inability to secure adequate prices in a particular country may not only limit the marketing of our products within that country, but may also adversely affect our ability to obtain acceptable prices in other markets. This may create the opportunity for third-party cross border trade or influence our decision to sell or not to sell a product, thus adversely affecting our geographic expansion plans and revenues.

The U.S. federal medical device excise tax may materially adversely affect our business and results of operations, and we may be subject to increased taxes in other jurisdictions.

The ACA imposed a 2.3% federal medical device excise tax on the sales in the United States of most medical devices. Most if not all of our products will be subject to this tax. This excise tax became effective in 2013 and has forced, and will continue to force us to identify ways to reduce spending in other areas to offset the expected earnings impact due to the tax. We do not expect to be able to pass along the cost of this tax to hospitals, which continue to face cuts to their Medicare reimbursement due to the ACA and the recently enacted ATRA. Nor do we expect to be able to offset the cost of the tax through higher sales volumes resulting from the expansion of health insurance coverage because of the demographics of the current uninsured population in the United States. While it is still too early to fully understand and predict the ultimate impact of the medical device tax on our business, ongoing implementation of this legislation and any similar taxes imposed in other jurisdictions could have a material adverse effect on our results of operations and cash flows. There are continuing efforts in the U.S. Congress to repeal this excise tax, but to date none of these efforts have been successful.

Risks Related to Our Intellectual Property and Litigation

If the combination of patents, trade secrets and contractual provisions that we rely on to protect our intellectual property is inadequate, our ability to commercialize our orthopedic products successfully will be harmed, and we may not be able to operate our business profitably.

Our success depends significantly on our ability to protect our proprietary rights to the technologies incorporated in our products. We currently have 34 issued U.S. patents, 38 pending U.S. patent applications, 11 granted foreign patents and 18 pending foreign patent applications. Our issued patents begin to expire in 2014, with the last of these patents expiring in 2031. We rely on a combination of patent protection, trade secret laws and nondisclosure, confidentiality and other contractual restrictions to protect our proprietary technology. However, these may not adequately protect our rights or permit us to gain or keep any competitive advantage.

The issuance of a patent is not conclusive as to its scope, validity or enforceability. The scope, validity or enforceability of our issued patents can be challenged in litigation or proceedings before the U.S. Patent and Trademark Office, or the USPTO, or foreign patent offices. In addition, our pending patent applications include claims to numerous important aspects of our products under development that are not currently protected by any of our issued patents. We cannot assure you that any of our pending patent applications will result in the issuance of patents to us. The USPTO or foreign patent offices may deny or require significant narrowing of claims in our pending patent applications. Patents issued as a result of the pending patent applications, if any, may not provide us with significant commercial protection or be issued in a form that is advantageous to us. Proceedings before the USPTO or foreign patent offices could result in adverse decisions as to the priority of our inventions and the narrowing or invalidation of claims in issued patents. The laws of some foreign countries may not protect our intellectual property rights to the same extent as the laws of the United States, if at all.

Our competitors may successfully challenge and invalidate or render unenforceable our issued patents, including any patents that may issue in the future, which could prevent or limit our ability to market our products and could limit our ability to stop competitors from marketing products that are substantially equivalent to ours. In addition, competitors may be able to design around our patents or develop products that provide outcomes that are comparable to our products but that are not covered by our patents.

CeramTec GmbH (“CeramTec”) has filed an opposition with the European Patent Office against our European Patent Number 2055267 (“EP ‘267 Patent”) entitled “Radiolucent bone graft,” which was granted by the European Patent Office. CeramTec’s Opposition includes allegations that the subject matter of EP ‘267 Patent lacks (i) adequate disclosure and (ii) novelty or at least an inventive step. The opposition is in the early stages, and if the European Patent Office rules that the allegations are correct in whole or in part, the claims of our European patent could be narrowed, or revoked in their entirety.

We have also entered into confidentiality and assignment of intellectual property agreements with all of our employees, consultants and advisors as one of the ways we seek to protect our intellectual property and other proprietary technology. However, these agreements may not be enforceable or may not provide meaningful protection for our trade secrets or other proprietary information in the event of unauthorized use or disclosure or other breaches of the agreements.

In the event a competitor infringes upon any of our patents or other intellectual property rights, enforcing our rights may be difficult, time consuming and expensive, and would divert management’s attention from managing our business. There can be no assurance that we will be successful on the merits in any enforcement effort. In addition, we may not have sufficient resources to litigate, enforce or defend our intellectual property rights.

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We have no patent protection covering the composition of matter for our solid *MC*² silicon nitride or the process we use for manufacturing our *MC*² silicon nitride, and competitors may create silicon nitride formulations substantially similar to ours.

Although we have a number of U.S. and foreign patents and pending applications relating to our *MC*² silicon nitride products or product candidates, we have no patent protection either for the composition of matter for our silicon nitride or for the processes of manufacturing *MC*² silicon nitride. As a result, competitors may create silicon nitride formulations substantially similar to ours, and use their formulations in products that may compete with our silicon nitride products, provided they do not violate our issued product patents. Although we have, and will continue to develop, significant know-how related to these processes, there can be no assurance that we will be able to maintain this know-how as trade secrets, and competitors may develop or acquire equally valuable or more valuable know-how related to the manufacture of silicon nitride.

We could become subject to intellectual property litigation that could be costly, result in the diversion of management's time and efforts, require us to pay damages, prevent us from marketing our commercially available products or product candidates and/or reduce the margins we may realize from our products that we may commercialize.

The medical devices industry is characterized by extensive litigation and administrative proceedings over patent and other intellectual property rights. Whether a product infringes a patent involves complex legal and factual issues, and the determination is often uncertain. There may be existing patents of which we are unaware that our products under development may inadvertently infringe. The likelihood that patent infringement claims may be brought against us increases as the number of participants in the orthopedic market increases and as we achieve more visibility in the market place and introduce products to market.

Any infringement claim against us, even if without merit, may cause us to incur substantial costs, and would place a significant strain on our financial resources, divert the attention of management from our core business, and harm our reputation. In some cases, litigation may be threatened or brought by a patent holding company or other adverse patent owner who has no relevant product revenues and against whom our patents may provide little or no deterrence. If we were found to infringe any patents, we could be required to pay substantial damages, including triple damages if an infringement is found to be willful, and royalties and could be prevented from selling our products unless we obtain a license or are able to redesign our products to avoid infringement. We may not be able to obtain a license enabling us to sell our products on reasonable terms, or at all, and there can be no assurance that we would be able to redesign our products in a way that would not infringe those patents. If we fail to obtain any required licenses or make any necessary changes to our technologies or the products that incorporate them, we may be unable to commercialize one or more of our products or may have to withdraw products from the market, all of which would have a material adverse effect on our business, financial condition and results of operations.

In addition, in order to further our product development efforts, we have entered into agreements with orthopedic surgeons to help us design and develop new products, and we expect to enter into similar agreements in the future. In certain instances, we have agreed to pay such surgeons royalties on sales of products which incorporate their product development contributions. There can be no assurance that surgeons with whom we have entered into such arrangements will not claim to be entitled to a royalty even if we do not believe that such products were developed by cooperative involvement between us and such surgeons. In addition, some of our surgeon advisors are employed by academic or medical institutions or have agreements with other orthopedic companies pursuant to which they have agreed to assign or are under an obligation to assign to those other companies or institutions their rights in inventions which they conceive or develop, or help conceive or develop.

There can be no assurance that one or more of these orthopedic companies or institutions will not claim ownership rights to an invention we develop in collaboration with our surgeon advisors or consultants on the basis that an agreement with such orthopedic company or institution gives it ownership rights in the invention or that our surgeon advisors or consultants otherwise have an obligation to assign such inventions to such company or institution. Any such claim against us, even without merit, may cause us to incur substantial costs, and would place a significant strain on our financial resources, divert the attention of management from our core business and harm our reputation.

We may be subject to damages resulting from claims that we, our employees, or our independent sales agencies have wrongfully used or disclosed alleged trade secrets of our competitors or are in breach of non-competition agreements with our competitors or non-solicitation agreements.

Many of our employees were previously employed at other orthopedic companies, including our competitors and potential competitors. Many of our distributors and potential distributors sell, or in the past have sold, products of our competitors. We may be subject to claims that either we, or these employees or distributors, have inadvertently or otherwise used or disclosed the trade secrets or other proprietary information of our competitors. In addition, we have been and may in the future be subject to claims that we

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caused an employee or sales agent to break the terms of his or her non-competition agreement or non-solicitation agreement. Litigation may be necessary to defend against these claims. Even if we are successful in defending against these claims, litigation could result in substantial costs and be a distraction to management. If we fail in defending such claims, in addition to paying money damages, we may lose valuable intellectual property rights or personnel. A loss of key personnel or their work product could hamper or prevent our ability to commercialize products, which could have an adverse effect on our business, financial condition and results of operations.

If our silicon nitride products or our product candidates conflict with the rights of others, we may not be able to manufacture or market our products or product candidates, which could have a material and adverse effect on us.

Our commercial success will depend in part on not infringing the patents or violating the other proprietary rights of third parties. Issued patents held by others may limit our ability to develop commercial products. All issued patents are entitled to a presumption of validity under the laws of the United States. If we need suitable licenses to such patents to permit us to develop or market our product candidates, we may be required to pay significant fees or royalties and we cannot be certain that we would even be able to obtain such licenses. Competitors or third parties may obtain patents that may cover subject matter we use in developing the technology required to bring our products to market, that we use in producing our products, or that we use in treating patients with our products. We know that others have filed patent applications in various jurisdictions that relate to several areas in which we are developing products. Some of these patent applications have already resulted in patents and some are still pending. If we were found to infringe any of these issued patents or any of the pending patent applications, when and if issued, we may be required to alter our processes or product candidates, pay licensing fees or cease activities. If use of technology incorporated into or used to produce our product candidates is challenged, or if our processes or product candidates conflict with patent rights of others, third parties could bring legal actions against us, in Europe, the United States and elsewhere, claiming damages and seeking to enjoin manufacturing and marketing of the affected products. Additionally, it is not possible to predict with certainty what patent claims may issue from pending applications. In the United States, for example, patent prosecution can proceed in secret prior to issuance of a patent, provided such application is not filed in foreign jurisdiction. For U.S. patent applications that are also filed in foreign jurisdictions, such patent applications will not publish until 18 months from the filing date of the application. As a result, third parties may be able to obtain patents with claims relating to our product candidates which they could attempt to assert against us. Further, as we develop our products, third parties may assert that we infringe the patents currently held or licensed by them, and we cannot predict the outcome of any such action.

There has been extensive litigation in the medical devices industry over patents and other proprietary rights. If we become involved in any litigation, it could consume a substantial portion of our resources, regardless of the outcome of the litigation. If these legal actions are successful, in addition to any potential liability for damages, we could be required to obtain a license, grant cross-licenses and pay substantial royalties in order to continue to manufacture or market the affected products.

We cannot assure you that we would prevail in any legal action or that any license required under a third party patent would be made available on acceptable terms, or at all. Ultimately, we could be prevented from commercializing a product, or forced to cease some aspect of our business operations, as a result of claims of patent infringement or violation of other intellectual property rights, which could have a material and adverse effect on our business, financial condition and results of operations.

Risks Related to Potential Litigation from Operating Our Business

We may become subject to potential product liability claims, and we may be required to pay damages that exceed our insurance coverage.

Our business exposes us to potential product liability claims that are inherent in the design, testing, manufacture, sale and distribution of our currently marketed products and each of our product candidates that we are seeking to introduce to the market. The use of orthopedic medical devices can involve significant risks of serious complications, including bleeding, nerve injury, paralysis, infection, and even death. Any product liability claim brought against us, with or without merit, could result in the increase of our product liability insurance rates or in our inability to secure coverage in the future on commercially reasonable terms, if at all. In addition, if our product liability insurance proves to be inadequate to pay a damage award, we may have to pay the excess of this award out of our cash reserves, which could significantly harm our financial condition. If longer-term patient results and experience indicate that our products or any component of a product causes tissue damage, motor impairment or other adverse effects, we could be subject to significant liability. A product liability claim, even one without merit, could harm our reputation in the industry, lead to significant legal fees, and result in the diversion of management's attention from managing our business.

Any claims relating to our improper handling, storage or disposal of biological or hazardous materials could be time consuming and costly.

Although we do not believe that the manufacture of our silicon nitride or non-silicon nitride products will involve the use of hazardous materials, it is possible that regulatory authorities may disagree or that changes to our manufacturing processes may result

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in such use. Our business and facilities and those of our suppliers and future suppliers may therefore be subject to foreign, federal, state and local laws and regulations governing the use, manufacture, storage, handling and disposal of hazardous materials and waste products. We may incur significant expenses in the future relating to any failure to comply with environmental laws. Any such future expenses or liability could have a significant negative impact on our business, financial condition and results of operations.

Risks Related to Our Common Stock

The price of our common stock is volatile and is likely to will continue to fluctuate due to reasons beyond our control.

The volatility of orthopedic company stocks often does not correlate to the operating performance of the companies represented by such stocks. Some of the factors that may cause the market price of our common stock to fluctuate include:

- our ability to sell our current products and the cost of revenue;
- our ability to develop, obtain regulatory clearances or approvals for, and market new and enhanced product candidates on a timely basis;
- changes in governmental regulations or in the status of our regulatory approvals, clearances or future applications;
- our announcements or our competitors' announcements regarding new products, product enhancements, significant contracts, number and productivity of distributors, number of hospitals and surgeons using products, acquisitions or strategic investments;
- announcements of technological or medical innovations for the treatment of orthopedic pathology;
- delays or other problems with the manufacturing of our products, product candidates and related instrumentation;
- volume and timing of orders for our products and our product candidates, if and when commercialized;
- changes in the availability of third-party reimbursement in the United States and other countries;
- quarterly variations in our or our competitors' results of operations;
- changes in earnings estimates or recommendations by securities analysts, if any, who cover our common stock;
- failure to meet estimates or recommendations by securities analysts, if any, who cover our stock;
- changes in the fair value of our common stock warrant liability resulting from changes in the market price of our common stock, which may result in significant fluctuations in our quarterly and annual operating results;
- changes in healthcare policy in the United States and internationally;
- product liability claims or other litigation involving us;
- sales of large blocks of our common stock, including sales by our executive officers, directors and significant stockholders;
- disputes or other developments with respect to intellectual property rights;
- changes in accounting principles;
- changes to tax policy; and
- general market conditions and other factors, including factors unrelated to our operating performance or the operating performance of our competitors.

These and other external factors may cause the market price and demand for our common stock to fluctuate substantially, which may limit or prevent our stockholders from readily selling their shares of common stock and may otherwise negatively affect the liquidity of our common stock. In addition, in the past, when the market price of a stock has been volatile, holders of that stock have sometimes instituted securities class action litigation against the company that issued the stock. If our stockholders brought a lawsuit against us, we could incur substantial costs defending the lawsuit regardless of the merits of the case or the eventual outcome. Such a lawsuit also would divert the time and attention of our management from running our company.

Securities analysts may not continue to provide coverage of our common stock or may issue negative reports, which may have a negative impact on the market price of our common stock.

Since completing our initial public offering of shares of our common stock in February 2014, a limited number of securities analysts have begun providing research coverage of our common stock. If securities analysts do not continue to cover our common stock, the lack of research coverage may cause the market price of our common stock to decline. The trading market for our common stock may be affected in part by the research and reports that industry or financial analysts publish about our business. If one or more of the analysts who elect to cover us downgrade our stock, our stock price would likely decline rapidly. If one or more of these

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analysts cease coverage of us, we could lose visibility in the market, which in turn could cause our stock price to decline. In addition, under the Sarbanes-Oxley Act of 2002, or the Sarbanes-Oxley Act, and a global settlement among the Securities and Exchange Commission, or the SEC, other regulatory agencies and a number of investment banks, which was reached in 2003, many investment banking firms are required to contract with independent financial analysts for their stock research. It may be difficult for a company such as ours, with a smaller market capitalization, to attract independent financial analysts that will cover our common stock. This could have a negative effect on the market price of our stock.

The low trading volume of our common stock may adversely affect the price of our shares.

Although our common stock is listed on the NASDAQ Capital Market, our common stock has experienced low trading volume. The average daily trading volume since our initial listing of our common stock on February 13, 2014 through March 14, 2014, as reported by NASDAQ, was approximately 219,000 shares. Limited trading volume may subject our common stock to greater price volatility and may make it difficult for investors to sell shares at a price that is attractive to them.

If our executive officers, directors and principal stockholders choose to act together, they will be able to exert significant influence over us and our significant corporate decisions and may act in a manner that advances their best interests and not necessarily those of other stockholders.

Our executive officers, directors, and beneficial owners of 5% or more of our outstanding common stock and their affiliates beneficially own approximately 13% of our outstanding common stock as of March 14, 2014. As a result, these persons, acting together, will have the ability to significantly influence the outcome of all matters requiring stockholder approval, including the election and removal of directors and any merger, consolidation, or sale of all or substantially all of our assets, and they may act in a manner that advances their best interests and not necessarily those of other stockholders, by among other things:

- delaying, deferring or preventing a change in control of our company;
- entrenching our management and/or our board of directors;
- impeding a merger, consolidation, takeover or other business combination involving our company;
- discouraging a potential acquirer from making a tender offer or otherwise attempting to obtain control of our company; or
- causing us to enter into transactions or agreements that are not in the best interests of all stockholders.

Future sales of our common stock in the public market may cause our stock price to decline and impair our ability to raise future capital through the sale of our equity securities.

There are a substantial number of shares of our common stock held by stockholders who owned shares of our capital stock prior to our initial public offering that will be able to be sold in the public market in the near future. Sales by such stockholders of a substantial number of shares could significantly reduce the market price of our common stock. Moreover, the holders of 2,581,941 shares of common stock, which were converted from convertible preferred stock to common stock upon the completion of our initial public offering, and holders of warrants to purchase 72,939 shares of common stock, which were converted from preferred stock warrants into common stock warrants upon the completion of our initial public offering, and holders of 12,363 shares of common stock, assuming the exercise of common stock warrants, will have rights, subject to some conditions, to require us to include their shares in registration statements that we may file for ourselves or other stockholders. These shares of common stock, totaling 2,667,243 shares, assuming the exercise of the common stock warrants, represent approximately 22% of the total number of shares of our common stock outstanding. In addition, warrants to acquire approximately 633,669 shares of our common stock and 8,627,454 shares of our outstanding common stock which are deemed to be "restricted securities" pursuant to Rule 144 under the Securities Act of 1933, as amended, or the Securities Act, will be eligible for sale in reliance on Rule 144, subject to the lock-up agreements covering substantially all of these securities which will expire on August 12, 2014. A holder of warrants to acquire shares of our common stock will be able to net exercise such warrants by surrendering a portion of that holder's warrants as payment of the exercise price rather than paying the exercise price in cash. As of December 31, 2013, holders of warrants to acquire approximately 633,669 shares of our common stock would be eligible to rely on Rule 144 for the resale of such shares if the warrants are net exercised, subject to the lock-up agreements. Additionally, all of our outstanding RSUs at March 6, 2014 will vest on August 12, 2014, resulting in approximately an additional 1,918,922 shares eligible to be sold in the public market.

We also intend to register all shares of our common stock that we may issue pursuant to our 2003 Stock Option Plan and our Amended and Restated 2012 Equity Incentive Plan, or the 2012 Plan in the second quarter of 2014. Shares issued by us upon exercise of options granted under our equity plans would be eligible for sale in the public market upon the effective date of the registration statement for those shares, subject to the lock-up agreements entered into in connection with the. If any of these holders cause a large number of securities to be sold in the public market, the sales could reduce the trading price of our common stock. These sales also could impede our ability to raise future capital.

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Anti-takeover provisions in our organizational documents and Delaware law may discourage or prevent a change in control, even if an acquisition would be beneficial to our stockholders, which could affect our stock price adversely and prevent attempts by our stockholders to replace or remove our current management.

Our restated certificate of incorporation and restated bylaws contain provisions that could discourage, delay or prevent a merger, acquisition or other change in control of our company or changes in our board of directors that our stockholders might consider favorable, including transactions in which you might receive a premium for your shares. These provisions also could limit the price that investors might be willing to pay in the future for shares of our common stock, thereby depressing the market price of our common stock. Stockholders who wish to participate in these transactions may not have the opportunity to do so. Furthermore, these provisions could prevent or frustrate attempts by our stockholders to replace or remove management. These provisions:

- allow the authorized number of directors to be changed only by resolution of our board of directors;
- provide for a classified board of directors, such that not all members of our board will be elected at one time;
- prohibit our stockholders from filling board vacancies, limit who may call stockholder meetings, and prohibit the taking of stockholder action by written consent;
- prohibit our stockholders from making certain changes to our restated certificate of incorporation or restated bylaws except with the approval of holders of 75% of the outstanding shares of our capital stock entitled to vote;
- require advance written notice of stockholder proposals that can be acted upon at stockholders meetings and of director nominations to our board of directors; and
- authorize our board of directors to create and issue, without prior stockholder approval, preferred stock that may have rights senior to those of our common stock and that, if issued, could operate as a “poison pill” to dilute the stock ownership of a potential hostile acquirer to prevent an acquisition that is not approved by our board of directors.

In addition, we are subject to the provisions of Section 203 of the Delaware General Corporation Law, which may prohibit certain business combinations with stockholders owning 15% or more of our outstanding voting stock. Any delay or prevention of a change in control transaction or changes in our board of directors could cause the market price of our common stock to decline.

We do not intend to pay cash dividends.

We have never declared or paid cash dividends on our capital stock and we do not anticipate paying any cash dividends in the foreseeable future. We currently intend to retain all available funds and any future earnings for debt service and use in the operation and expansion of our business. The GE Secured Lending Facility also contains a negative covenant which prohibits us from paying dividends to our stockholders without the prior written consent of GE Capital. In addition, the terms of any future debt or credit facility may preclude us from paying any dividends.

Risks Related to Public Companies

We are an “emerging growth company” as defined in the Jumpstart Our Business Startups Act of 2012 and a “smaller reporting company” and the reduced disclosure requirements applicable to emerging growth companies and smaller reporting companies may make our common stock less attractive to investors.

We are an “emerging growth company,” as defined in the Jumpstart Our Business Startups Act of 2012, or the JOBS Act. For as long as we continue to be an emerging growth company, we may take advantage of exemptions from various reporting requirements that are applicable to other public companies that are not emerging growth companies, including (1) not being required to comply with the auditor attestation requirements of Section 404 of the Sarbanes-Oxley Act, (2) reduced disclosure obligations regarding executive compensation in our periodic reports and proxy statements and (3) exemptions from the requirements of holding a nonbinding advisory vote on executive compensation and stockholder approval of any golden parachute payments not previously approved. Additionally, under the JOBS Act, emerging growth companies can also delay adopting new or revised accounting standards until such time as those standards apply to private companies. We are electing to delay such adoption of new or revised accounting standards on the relevant dates on which adoption of such standards is required for non-emerging growth companies. As a result of this election, our financial statements may not be comparable to the financial statements of other public companies.

We may take advantage of these exemptions until we are no longer an emerging growth company. Under the JOBS Act, we may be able to maintain emerging growth company status for up to five years, although circumstances could cause us to lose that status earlier, including if the market value of our common stock held by non-affiliates exceeds \$700 million as of any June 30 before the end of such five-year period or if we have total annual gross revenue of \$1.0 billion or more during any fiscal year before that time, in

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which cases we would no longer be an emerging growth company as of the following December 31. Additionally, if we issue more than \$1.0 billion in non-convertible debt during any three-year period before that time, we would cease to be an emerging growth company immediately.

We are also currently a “smaller reporting company” as defined in the Securities Exchange Act of 1934, and in the event that we are still considered a smaller reporting company at such time as we cease being an emerging growth company, we will be required to provide additional disclosure in our SEC filings. However, similar to emerging growth companies, smaller reporting companies are able to provide simplified executive compensation disclosures in their filings, are exempt from the provisions of Section 404(b) of the Sarbanes-Oxley Act requiring that independent registered public accounting firms provide an attestation report on the effectiveness of internal control over financial reporting, and have certain other decreased disclosure obligations in their SEC filings, including, among other things, only being required to provide two years of audited financial statements in annual reports. We cannot predict whether investors will find our common stock less attractive because of our reliance on any of these exemptions. If some investors find our common stock less attractive as a result, there may be a less active trading market for our common stock and our stock price may be more volatile.

We incur substantial costs as a result of being a public company and our management expects to devote substantial time to public company compliance programs.

As a public company, we incur significant legal, insurance, accounting and other expenses, including costs associated with public company reporting. We intend to invest resources to comply with evolving laws, regulations and standards, and this investment will result in increased general and administrative expenses and may divert management’s time and attention from product development and commercialization activities. If our efforts to comply with new laws, regulations and standards differ from the activities intended by regulatory or governing bodies due to ambiguities related to practice, regulatory authorities may initiate legal proceedings against us, and our business may be harmed. These laws and regulations could make it more difficult and costly for us to obtain director and officer liability insurance for our directors and officers, and we may be required to accept reduced coverage or incur substantially higher costs to obtain coverage. These factors could also make it more difficult for us to attract and retain qualified executive officers and qualified members of our board of directors, particularly to serve on our audit and compensation committees. In addition, if we are unable to continue to meet the legal, regulatory and other requirements related to being a public company, we may not be able to maintain the listing of our common stock on The NASDAQ Capital Market, which would likely have a material adverse effect on the trading price of our common stock.

Our internal control over financial reporting does not currently meet the standards required by Section 404 of the Sarbanes-Oxley Act, and failure to achieve and maintain effective internal control over financial reporting in accordance with Section 404 of the Sarbanes-Oxley Act could result in material misstatements of our annual or interim financial statements and have a material adverse effect on our business and share price.

As a new public company, we are not currently required to make a formal assessment of the effectiveness of our internal control over financial reporting for purposes of compliance with the SEC’s rules that implement Section 404 of the Sarbanes Oxley Act. We are, however, required to comply with certain of these rules, which will require management to certify financial and other information in our quarterly and annual reports and provide an annual management report on the effectiveness of our internal control over financial reporting commencing with our second annual report. This assessment will need to include the disclosure of any material weaknesses or significant deficiencies in our internal control over financial reporting identified by our management or our independent registered public accounting firm. A “material weakness” is a deficiency, or a combination of deficiencies, in internal control over financial reporting such that there is a reasonable possibility that a material misstatement of our annual or interim financial statements will not be prevented or detected on a timely basis. A “significant deficiency” is a deficiency, or a combination of deficiencies, in internal control over financial reporting that is less severe than a material weakness, yet important enough to merit attention by those responsible for oversight of our financial reporting, including the audit committee of the board of directors.

Our independent registered public accounting firm will not be required to formally attest to the effectiveness of our internal control over financial reporting until the later of our second annual report or the first annual report required to be filed with the SEC following the date we are no longer an “emerging growth company” as defined in the JOBS Act. However, in connection with our audits for the years ended December 31, 2013 and 2012, and their review of our interim financial statements, our independent registered public accounting firm noted four material weaknesses and one significant deficiency in our internal control over financial reporting.

One material weakness related to our improper recording and disclosure of non-routine transactions due to deficiencies in the design and operation of our controls to account for non-routine transactions as part of the financial close process. We plan to remedy this by increasing the size and expertise of our internal accounting team.

Another material weakness was identified related to the deficiency in the design and operation of our controls to account for inventory. In addition to increasing the size and expertise of our accounting team, we plan to address this deficiency by physically counting inventory held by certain of our distributors on a more frequent basis and monitoring more closely the movement of inventory between locations.

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The third material weakness related to deficiencies in our income tax accounting. We intend to implement a formal process for accounting for income taxes, including evaluating the tax treatment of certain transactions on permanent and temporary book/tax differences, and the effect on the income tax provision and related deferred tax accounting balances.

The fourth material weakness relates to deficiencies in the design and operation of our controls to appropriately identify and evaluate transactions for appropriate cut-off at the end of the financial reporting period and the level of precision and timeliness of our financial close process. In addition to increasing the size and expertise of our accounting team, we plan to remedy this by implementing a formal financial close process related to financial reporting.

Additionally, our independent registered public accounting firm identified a significant deficiency related to the design and operation of our controls to manage the safeguarding of assets, particularly our instruments that we provide to surgeons and hospitals on consignment. We plan to implement a formal process for tracking and monitoring fixed assets as they are deployed for use at various locations.

We cannot assure you that our plans will sufficiently address the identified deficiencies, nor can we assure you that there will not be material weaknesses or significant deficiencies in our internal controls in the future. Additionally, in the event that our internal control over financial reporting is perceived as inadequate, or that we are unable to produce timely or accurate financial statements, investors may lose confidence in our operating results and the trading price of our common stock could decline.

Finally, as a private company, we were not previously required to prepare quarterly financial statements, nor were we required to generate financial statements in the time periods mandated for public companies by the SEC's reporting requirements. We believe that we will need to expand our accounting resources, including the size and expertise of our internal accounting team, to effectively execute a quarterly close process and on an appropriate time frame for a public company. If we are unsuccessful or unable to sufficiently expand these resources, we may not be able to produce U.S. GAAP-compliant financial statements on a time frame required to comply with our reporting requirements under the Exchange Act, and the financial statements we produce may contain material misstatements, either of which could cause investors to lose confidence in our financial reports and our financial reporting generally, which could lead to a material decline in the trading price of our common stock.

ITEM 1B. UNRESOLVED STAFF COMMENTS

Not applicable.

ITEM 2. PROPERTIES

Our 54,000 square foot corporate office and manufacturing facilities are located in Salt Lake City, Utah. We occupy these facilities pursuant to a lease that expires in January 2020. Pursuant to the terms of the lease agreement, we may extend the lease for two additional periods of five years each. We believe that our existing facilities are adequate for our current and projected needs for the foreseeable future.

ITEM 3. LEGAL PROCEEDINGS

We are currently not a party to any material legal proceedings. However, our industry is characterized by frequent claims and litigation, including claims regarding intellectual property and product liability. As a result, we may be subject to various legal proceedings in the future.

ITEM 4. MINE SAFETY DISCLOSURES

This item does not apply to our business.

PART II

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

Market Information

Our common stock began trading on The NASDAQ Stock Market on February 13, 2014 under the symbol "AMDA." At December 31, 2013, our common stock was privately held and there was no established public trading market for our common stock. As a result, we have not set forth quarterly information with respect to the high and low prices for our common stock for the two most recent fiscal years. Unless otherwise indicated, all information contained in this Annual Report reflects a 1-for-25.7746 reverse split of our common stock, which was effected on February 11, 2014.

Holdings of Record

As of March 14, 2014, we had approximately 597 holders of record of our common stock. Because many of our shares of common stock are held by brokers and other institutions on behalf of stockholders, this number is not indicative of the total number of stockholders represented by these stockholders of record.

Dividends

We have not paid dividends to stockholders since inception and do not plan to pay cash dividends in the foreseeable future. We currently intend to retain earnings, if any, to finance our growth.

Unregistered Sales of Securities

a) Issuances of Capital Stock and Warrants

The sale and issuance of the securities set forth below were deemed to be exempt from registration under the Securities Act by virtue of Section 4(2) or Rule 506 promulgated under Regulation D promulgated thereunder and Section 3(a)(9). Each of the recipients of securities in these transactions was an accredited investor within the meaning of Rule 501 of Regulation D under the Securities Act and had adequate access, through employment, business or other relationships, to information about us. No underwriters were involved in these transactions.

- In March 2013, we issued an aggregate of 178,516 shares of our common stock to 33 accredited investors upon exercise of warrants and the sale of additional shares of our common stock to other investors at \$17.53 per share for an aggregate purchase price of \$3,128,802. We also issued each investor purchasing shares of our common stock through the exercise of warrants new warrants to purchase an aggregate of 76,455 shares of our common stock at an exercise price of \$17.53 per share.
- In 2013, we issued a total of 13,191 shares of our common stock at \$17.53 per share to sales agents for equity incentive awards earned in 2011, 2012 and 2013 in connection with our obligations under a settlement agreement dated May 1, 2010.
- On August 30, 2013 and September 20, 2013, we issued and sold a total of 94.8 units, each unit consisting of 50,000 shares of our Series F convertible preferred stock and a warrant to acquire 970 shares of our common stock at an exercise price of \$25.77 per share, to 45 accredited investors at \$100,000 per unit for an aggregate purchase price of \$9,480,000. The purchase of these units resulted in our issuance of 4,740,000 shares of our Series F convertible preferred stock and warrants to purchase 91,951 shares of our common stock. In connection with this offering, we issued warrants to purchase an aggregate of 9,311 shares of our common stock, at an exercise price of \$56.70 per share, to TGP Securities, Inc.

(b) Certain Equity Grants and Exercises of Stock Options

In 2013, we granted 13,191 options to purchase shares of our common stock and granted a total of 197,335 RSUs. During the same period, 60,718 options to purchase shares of common stock were exercised.

Option grants, RSU grants and the issuances of common stock upon exercise of such options were exempt pursuant to Rule 701 and Section 4(2) of the Securities Act of 1933.

Use of Proceeds from Sale of Registered Securities

On February 12, 2014, our registration statement on Form S-1 (File No. 333-192232) was declared effective for our initial public offering of shares of our common stock, or the IPO, pursuant to which we registered the sale of 3,500,000 shares of our common stock

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at \$5.75 per share, plus up to an additional 525,000 additional shares to cover the underwriters' overallotment option. The IPO generated net proceeds to us of approximately \$15.4 million, after deducting underwriting discounts and expenses of approximately \$5.8 million. The IPO commenced as of February 12, 2014, and terminated before all of the securities registered under the registration statements were sold because the underwriters elected to partially exercise their overallotment option by purchasing 182,900 additional shares of our common stock. JMP Securities LLC acted as the sole book-running manager for the offering. Needham & Company, LLC acted as co-manager for the IPO.

There has been no material change in the planned use of proceeds from the IPO as described in our final prospectus filed with the SEC on February 12, 2014 pursuant to Rule 424(b).

Issuer Purchases of Equity Securities

None

ITEM 6. SELECTED FINANCIAL DATA

Not applicable.

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

You should read the following discussion and analysis of our financial condition and results of operations together with our consolidated financial statements and related notes appearing elsewhere in this Annual Report. This discussion and analysis contains forward-looking statements based upon current beliefs, plans, expectations, intentions and projections that involve risks, uncertainties and assumptions, such as statements regarding our plans, objectives, expectations, intentions and projections. Our actual results and the timing of selected events could differ materially from those anticipated in these forward-looking statements as a result of several factors, including those set forth under "Risk Factors" and elsewhere in this Annual Report.

Overview

We are a commercial biomaterial company focused on using our silicon nitride technology platform to develop, manufacture and sell a broad range of medical devices. We currently market spinal fusion products and are developing products for use in total hip and knee joint replacements. We believe our silicon nitride technology platform enables us to offer new and transformative products in the orthopedic and other medical device markets. We believe we are the first and only company to use silicon nitride in medical applications and over 14,000 of our intervertebral fusion devices have been implanted in patients.

We currently market our *Valeo MC²* silicon nitride interbody spinal fusion devices in the United States and Europe for use in the cervical and thoracolumbar areas of the spine. We believe our *Valeo* devices have a number of advantages over existing products due to silicon nitride's key characteristics, resulting in faster and more effective fusion and reduced risk of infection. Our first generation *Valeo* silicon nitride device received 510(k) regulatory clearance and a CE Mark in 2008. Based on surgeon feedback, we developed a second generation of *Valeo* AL, PL and OL products with design enhancements that improve surgeon control during implantation and stability post procedure. In 2013, we initiated a targeted launch of our second generation *Valeo* AL, PL and OL interbody fusion devices and expect to complete the full launch in the second quarter of 2014. We are also completing the development of our second generation Cervical and TL *Valeo* interbody spinal fusion device and expect these to be launched in the second half of 2014. We also market our *Valeo* composite interbody spinal fusion device made from both our solid *MC²* and porous *C⁵C* silicon nitride in the Netherlands, Spain and Germany. We are currently conducting a prospective clinical trial in Europe, named CASCADE, comparing our *Valeo* composite silicon nitride interbody devices to PEEK interbody devices to obtain additional data to support 510(k) clearance in the United States. The trial is 100% enrolled. We expect results to be available in the second half of 2014. If this trial is successful, we plan to file a 510(k) submission with the FDA by mid-2015. In addition, in the first half of 2013, we initiated a Design and Build Program focused on collaborating with influential surgeons to develop customized silicon nitride spinal fusion products and instruments and the first products designed under this program were sold in the third quarter of 2013.

In addition to our silicon nitride-based spinal fusion products, we market a complementary line of non-silicon nitride spinal fusion products which allows us to provide surgeons and hospitals with a broader range of products. These products include three lines of spinal fusion devices and five types of orthobiologics, which are used by surgeons to help promote bone growth and fusion in spinal fusion procedures. Although our non-silicon nitride products have accounted for approximately 66% and 74% or more of our product revenues for the years ended December 31, 2013 and 2012, respectively, we believe the continued promotion and potential for adoption of our silicon nitride products and product candidates, if approved, provides us the greatest opportunity to grow our business in new and existing markets and achieve our goal to become a leading biomaterial company.

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We market and sell our products to surgeons and hospitals in the United States and select markets in Europe and South America through our established network of more than 50 independent sales distributors. A substantial portion of our product revenue has historically been derived from sales in the United States. Our largest customer, Bon Secours St. Mary's Hospital, accounted for 14% of our product revenues for each of the years ended December 31, 2013 and 2012. A significant portion of this hospital group's purchases from us are non-silicon nitride products and its accounts receivable balance was approximately 15% of our total trade accounts receivable at December 31, 2013.

We plan to use our silicon nitride technology platform to expand our product offerings. We are incorporating our silicon nitride technology into components for use in total hip and knee replacement product candidates that we are, or plan on, developing in collaboration with a strategic partner. In addition, we believe our silicon nitride technology platform can be used for developing products in other markets and have developed prototypes for use in the dental, sports medicine and trauma markets. We believe our coating technology may be used to enhance our metal products as well as commercially-available metals, such as those used in spinal fusion, joint replacement and other medical products.

Components of our Results of Operations

We manage our business within one reportable segment, which is consistent with how our management reviews our business, makes investment and resource allocation decisions and assesses operating performance.

Product Revenue

We derive our product revenue primarily from the sale of spinal fusion devices and related products used in the treatment of spine disorders. Our product revenue is generated from sales to two types of customers: (1) surgeons and hospitals; and (2) stocking distributors. Most of our products are sold on a consignment basis through a network of independent sales distributors; however, we also sell our products to independent stocking distributors. Product revenue is recognized when all four of the following criteria are met: (1) persuasive evidence that an arrangement exists; (2) delivery of the products has occurred; (3) the selling price of the product is fixed or determinable; and (4) collectability is reasonably assured. We generate the majority of our revenue from the sale of inventory that is consigned to independent sales distributors that sell our products to surgeons and hospitals. For these products, we recognize revenue at the time we are notified the product has been used or implanted and a valid purchase order has been received. For all other transactions, we recognize revenue when title and risk of loss transfer to the stocking distributor, and all other revenue recognition criteria have been met. We generally recognize revenue from sales to stocking distributors at the time the product is shipped to the distributor. Stocking distributors, who sell the products to their customers, take title to the products and assume all risks of ownership at time of shipment. Our stocking distributors are obligated to pay within specified terms regardless of when, if ever, they sell the products. Our policy is to classify shipping and handling costs billed to customers as an offset to total shipping expense in the statement of operations, primarily within sales and marketing. In general, our customers do not have any rights of return or exchange.

We believe our product revenue from the sale of our silicon nitride based products and our non-silicon nitride products will increase due to our sales and marketing efforts and as we introduce new silicon nitride based products into the market, such as our second generation *Valeo* interbody spinal fusion products in the United States. We expect that our product revenue will continue to be primarily attributable to sales of our products in the United States, though, as we expand our sales and marketing efforts and market additional products abroad, such as our spinal fusion device incorporating our *C^{SC}*, we expect international sales will increase.

Cost of Revenue

The expenses that are included in cost of revenue include all direct product costs if we obtained the product from third-party manufacturers and our in-house manufacturing costs for the products we manufacture. We obtain our non-silicon nitride products, including our metal and orthobiologic products, from third-party manufacturers, while we manufacture our silicon-nitride products in-house.

Specific provisions for excess or obsolete inventory and, beginning in 2013, the 2.3% excise tax on the sale of medical devices in the United States, are also included in cost of revenue. In addition, we pay royalties attributable to the sale of specific products to some of our surgeon advisors that assisted us in the design, clearance or commercialization of a particular product, and these payments are recorded as cost of revenue.

Gross Profit

Our gross profit measures our product revenue relative to our cost of revenue. While we expect our cost of revenue to increase in absolute terms as our sales volume increases, we believe our gross profit will be higher as we realize manufacturing efficiencies associated with our silicon nitride-based products.

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Research and Development Expenses

Our net research and development costs are expensed as incurred. Research and development costs consist of engineering, product development, clinical trials, test-part manufacturing, testing, developing and validating the manufacturing process, manufacturing, facility and regulatory-related costs. Research and development expenses also include employee compensation, employee and non-employee stock-based compensation, supplies and materials, consultant services, and travel and facilities expenses related to research activities. To the extent that certain research and development expenses are directly related to our manufactured products, such expenses and related overhead costs are allocated to inventory.

We expect to incur additional research and development costs as we continue to develop new spinal fusion products such as our second generation *Valeo* products, our product candidates for total joint replacements, such as our total hip replacement product candidate, and our silicon nitride-coated metals which may increase our research and development expenses.

Sales and Marketing Expenses

Sales and marketing expenses consist of salaries, benefits and other related costs, including stock-based compensation, for personnel employed in sales, marketing, medical education and training. In addition, our sales and marketing expenses include commissions and bonuses, generally based on a percentage of sales, to our sales managers and independent sales distributors. We provide our products in kits or banks that consist of a range of device sizes and separate instruments necessary to complete the surgical procedure. We generally consign our instruments to our distributors or our hospital customers that purchase the device used in spinal fusion surgery. Our sales and marketing expenses include depreciation of the surgical instruments.

We expect our sales and marketing expenses to continue to increase, including instrument set depreciation, as we introduce new products, such as our second generation *Valeo* spinal fusion products into the United States, and seek to enhance our commercial infrastructure, including increasing our marketing efforts and further educating our distributors. Additionally, we expect our commissions to continue to increase in absolute terms over time but remain approximately the same or decrease as a percentage of product revenue.

General and Administrative Expenses

General and administrative expenses primarily consist of salaries, benefits and other related costs, including stock-based compensation, for certain members of our executive team and other personnel employed in finance, legal, compliance, administrative, information technology, customer service, executive and human resource departments. General and administrative expenses include allocated facility expenses, related travel expenses and professional fees for accounting and legal services.

We expect our general and administrative expenses will increase due to costs associated with transitioning from a private to a public company and as we continue to grow our business.

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Results of Operations

Year Ended December 31, 2013 Compared to the Year Ended December 31, 2012

The following table sets forth, for the periods indicated, our results of operations for the years ended December 31, 2013 and 2012 (in thousands):

	Year Ended December 31,		\$ Change	% Change
	2013	2012		
Product revenue	\$ 22,314	\$ 23,065	\$ (751)	(3%)
Total cost of revenue	7,045	6,466	579	9%
Gross profit	15,269	16,599	(1,330)	(8%)
Operating expenses:				
Research and development	3,461	6,013	(2,552)	(42%)
General and administrative	5,759	7,313	(1,554)	(21%)
Sales and marketing	16,384	17,094	(710)	(4%)
Impairment loss on intangible assets	—	15,281	(15,281)	N/A
Total operating expenses	25,604	45,701	(20,097)	(44%)
Loss from operations	(10,335)	(29,102)	18,767	(64%)
Other income (expense), net	2,049	(6,659)	8,708	131%
Net loss before income taxes	(8,286)	(35,761)	27,475	77%
Income tax benefit	—	726	726	N/A
Net loss	\$ (8,286)	\$ (35,035)	\$ 26,749	76%

Product Revenue

The following table sets forth our product revenue from sales of the indicated product category for the years ended December 31, 2013 and 2012 (in thousands):

	Year Ended December 31,		\$ Change	% Change
	2013	2012		
Silicon Nitride	\$ 7,667	\$ 6,578	\$ 1,089	17%
Non-Silicon Nitride	14,647	16,487	(1,840)	(11%)
Total Product Revenue	\$ 22,314	\$ 23,065	\$ (751)	(3%)

Total product revenue was \$22.3 million in 2013 as compared to \$23.1 million in 2012, a decrease of \$0.8 million or 3%. This decrease in total product revenue was primarily attributable to our restructuring of our sales and marketing teams during the first quarter of 2013, resulting from changes in our distribution network, the timing of the launch of our second generation *Valeo* products and a one-time sale of non-silicon nitride products to an international customer in the 2012 period with no corresponding sale in 2013. Sales of our silicon nitride products increased by \$1.1 million, or 17%, in 2013 as compared to 2012. Non-silicon nitride sales decreased \$1.8 million, or 11%, in 2013 compared to 2012, as the new sales team was primarily focused on silicon nitride product sales as opposed to non-silicon nitride product sales.

The following table sets forth, for the periods indicated, our product revenue by geographic area (in thousands):

	Year Ended December 31,		\$ Change	% Change
	2013	2012		
Domestic	\$ 22,203	\$ 21,847	\$ 356	2%
International	111	1,218	(1,107)	(91%)
Total Product Revenue	\$ 22,314	\$ 23,065	\$ (751)	(3%)

Product revenue attributable to sales in the United States was \$22.2 million in 2013, an increase of \$0.4 million, or 2%, as compared to 2012. Product revenue attributable to international sales was \$0.1 million in 2013, a decrease of \$1.1 million, or 91%, as compared to 2012. The decrease was primarily attributable to a one-time sale of non-silicon nitride products to an international customer in 2012.

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Cost of Revenue

Cost of revenue was \$7.0 million in 2013 as compared to \$6.5 million in 2012, an increase of \$0.5 million, or 8%. This increase was primarily related to an increase in excess and obsolete inventory costs of \$0.3 million related to our first generation *Valeo* products and the new 2.3% medical device excise tax in the United States, which totaled \$0.4 million in 2013. The increases were partially offset by lower costs of our orthobiologic products in 2013 as compared to 2012.

Gross Profit

Gross profit as a percentage of product revenue decreased by 4% to 68% for 2013 from 72% for the same period in 2012, primarily as a result of the U.S. medical device excise tax of 2.3% on product revenue which became effective in January 2013 and an increase in excess and obsolete inventory costs of \$0.3 million related to our first generation *Valeo* products.

Research and Development Expenses

Research and development expenses were \$3.5 million in 2013 as compared to \$6.0 million in 2012, a decrease of \$2.5 million, or 42%. This decrease was primarily due to our allocation, in 2013, of an additional \$2.9 million of overhead costs to inventory as a result of the ramp-up phase for our second generation *Valeo* products, which overhead costs had been allocated to research and development expenses in 2012.

General and Administrative Expenses

General and administrative expenses were \$5.8 million in 2013 as compared to \$7.3 million in 2012, a decrease of \$1.5 million, or 21%. This decrease was primarily due to decreases of \$1.4 million in amortization expense and \$1.1 million in legal and patent expense, partially offset by a \$0.8 million increase in employee compensation and \$0.2 million increase in accounting and consulting expenses.

Sales and Marketing Expenses

Sales and marketing expenses were \$16.4 million in 2013 as compared to \$17.1 million in 2012, a decrease of \$0.7 million, or 4%. This decrease was primarily due to a decrease of \$0.8 million in marketing expenses and \$0.5 million decrease in depreciation and maintenance expense associated with instrumentations. These decreases were partially offset by an increase of \$0.6 million in commission expenses, primarily to our sales distributors, to support increased silicon nitride sales volume.

Impairment Loss on Intangible Assets

Impairment loss on intangible assets was \$15.3 million in 2012 relating to assets we obtained in our acquisition of US Spine, Inc. ("US Spine"). The amount of the impairment loss was determined during management's annual impairment review and resulted from lower sales of certain products and lower sales to customers acquired in the US Spine transaction than originally expected. There was not an impairment loss on intangible assets during 2013.

Other Income (Expense), Net

Other income was \$2.0 million in 2013 as compared to other expenses of \$6.7 million in 2012, a decrease of \$8.7 million, or 131%. This decrease in other expense was primarily due to a net decrease in fair value of \$4.7 million of our common and preferred stock warrant liabilities as a result of our lower stock price at December 31, 2013 as compared to December 31, 2012. Furthermore, there was a decrease of \$3.8 million in interest expense due to our senior secured notes being converted into shares of Series F convertible preferred stock in December 2012.

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Liquidity and Capital Resources

For the years ended December 31, 2013 and 2012, we incurred a net loss of \$8.3 million and \$35.0 million, respectively, and used cash in operations of \$9.9 million and \$9.7 million, respectively. We have an accumulated deficit of \$139.9 million and \$131.6 million as of December 31, 2013 and 2012, respectively. With the exception of a small net income for the years ended December 31, 2002 and 1999, we have incurred net losses in each year since inception. To date, our operations have been principally financed from proceeds from the issuance of convertible preferred stock and common stock, convertible debt and bank debt and, to a lesser extent, cash generated from product sales. Since January 2011 and up to December 31, 2013, we issued the following securities to help fund our operations:

- between March 2011 and February 2012, we issued aggregate principal amount of \$29.8 million of Senior Secured Subordinated 6%/8% Convertible Promissory Notes, or the Senior Secured Notes, and warrants to purchase an aggregate of 288,685 shares of our common stock at an exercise price of \$51.55 per share. All outstanding Senior Secured Notes were converted into 14,887,500 shares of our Series F convertible preferred stock in December 2012 contemporaneously with our entering into a new term loan and a revolving credit facility with General Electric Capital Corporation, or GE Capital, and Zions First National Bank, or the GE Secured Lending Facility;
- in February 2013, we issued an aggregate of 178,516 shares of our common stock upon exercise of warrants and the sale of additional shares of our common stock at \$17.53 per share for an aggregate purchase price of \$3.1 million. We also issued each investor purchasing shares of our common stock through the exercise of warrants new warrants to purchase shares of our common stock at an exercise price of \$17.53 per share; and
- in August and September 2013, we issued an aggregate of 94.8 units, each unit consisting of 50,000 shares of our Series F convertible preferred stock and a warrant to acquire 970 shares of our common stock at an exercise price of \$25.77 per share, for gross proceeds of \$9.5 million.

As of December 31, 2013 and 2012, we had approximately \$2.7 million and \$5.7 million, respectively, in cash, cash equivalents, restricted cash and marketable securities. Restricted cash, which was \$0.4 million and \$0.3 million at December 31, 2013 and 2012, respectively, consists of cash balances in transit from a segregated account that must first be applied to pay down any outstanding balance on the revolving credit facility portion of the GE Secured Lending Facility. In order to finance the continued growth in product sales, to invest in further product development and to otherwise satisfy obligations as they mature, we completed an initial public offering of our common stock ("IPO"), in which we sold and issued 3,682,900 shares, including 182,900 shares sold pursuant to the exercise by the underwriters of their over-allotment option, in February 2014, at an issuance price of \$5.75 per share, less underwriting discounts and commissions. As a result of the IPO, we received proceeds of approximately \$15.4 million, net of approximately \$5.8 million in IPO related costs. We will need to obtain additional funding during the third quarter of 2014 to satisfy the \$7.2 million of principal obligations due under the GE Secured Lending Facility in 2014 (which amount excludes interest payments and any other fees and charges incurred or that we may incur) and to maintain compliance with the financial and liquidity covenants related to the GE Secured Lending Facility through 2014. Furthermore, if we are unable to access additional funds prior to becoming non-compliant with the financial or liquidity covenants, the entire remaining balance of the GE Secured Lending Facility could become immediately due and payable at the option of GE Capital.

In addition, the repayment of the GE Secured Lending Facility and the liquidity covenant limit our ability to use our cash and cash equivalents to fund our operations and may restrict our ability to continue development of our product candidates. Additionally, the GE Secured Lending Facility restricts our ability to incur additional pari passu indebtedness, which may reduce our ability to seek additional financing. If adequate funds are not available on a timely basis, we may terminate or delay the development of one or more of our product candidates, or delay activities necessary to commercialize our product candidates. Additional funding may not be available to us on acceptable terms, or at all. Any additional equity financing, if available, may not be available on favorable terms and will most likely be dilutive to our current stockholders, and debt financing, if available, may involve more restrictive covenants. Our ability to access capital when needed is not assured and, if not achieved on a timely basis, will materially harm our business, financial condition and results of operations.

We are seeking additional financing through the issuance of shares of our common stock and/or obtaining additional debt financing, including a refinancing of our existing debt, to satisfy our debt obligations, meet our working capital requirements, make continued investment in research and development and make capital expenditures needed for us to maintain and expand our business. We may not be able to obtain additional financing on terms favorable to us, if at all. It is also possible that we may allocate significant amounts of capital toward solutions or technologies for which market demand is lower than anticipated and, as a result, abandon such efforts. If we are unable to obtain adequate financing or financing on terms satisfactory to us when we require it, or if we expend capital on projects that are not successful, our ability to continue to support our business growth and to respond to business challenges could be significantly limited, or we may even have to scale back our operations. If we raise additional funds through further issuances of equity or convertible debt securities, our existing stockholders could suffer significant dilution, and any new equity securities we issue could have rights, preferences and privileges superior to those of holders of our common stock.

Pursuant to its terms, we must repay our \$18.0 million term loan with GE Capital by making principal payments of \$600,000 over a period of 30 months, which began in January 2014. We must pay GE Capital a repayment fee of \$720,000 upon prepayment in full or at scheduled maturity of the term loan. The GE Secured Lending Facility also has minimum liquidity covenants that require us to maintain minimum levels of cash, cash equivalents and availability under the revolving credit facility, which can restrict our ability to use our cash and cash equivalents. We were in default of this liquidity covenant in November 2013, and, in December 2013, we amended the terms of the GE Secured Lending Facility to allow for a temporary waiver, effective from November 1, 2013 through January 31, 2014, of the liquidity covenant under the agreement for a fee of \$860,000, payable in March 2014. In addition, we agreed to an additional credit reserve in the amount of \$0.5 million, bringing the total reserve to \$1.0 million. On January 28, 2014, we obtained an additional waiver of the liquidity covenant from GE Capital through February 28, 2014 and agreed to increase the credit reserve under this facility by an additional \$0.5 million, bringing the total reserve to \$1.5 million. The total accrued amendment fees at December 31, 2013 were \$1.1 million. We also

agreed to pay GE Capital a fee of \$200,000 on March 31, 2014 in connection with

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the January 28, 2014 waiver if the GE Secured Lending Facility is not repaid on or before March 31, 2014. Accordingly, we expect to pay the \$1.3 million in amendment fees in April 2014. Although we were in compliance with the liquidity covenant at February 28, 2014, we anticipate that we will be unable to comply with the liquidity covenant prior to December 31, 2014, and have therefore classified the entire obligation as a current liability.

Going Concern

Our ability to access capital when needed is not assured and, if not achieved on a timely basis, will materially harm our business, financial condition and results of operations. These uncertainties create substantial doubt about our ability to continue as a going concern. Our independent registered public accounting firm included an explanatory paragraph regarding substantial doubt about our ability to continue as a going concern in their report on our annual financial statements for the fiscal year ended December 31, 2013. The financial information throughout this Annual Report have been prepared on a basis which assumes that we will continue as a going concern, which contemplates the realization of assets and the satisfaction of liabilities and commitments in the normal course of business. This financial information and statements do not include any adjustments that may result from the outcome of this uncertainty.

Cash Flows

The following table summarizes, for the periods indicated, cash flows from operating, investing and financing activities (in thousands):

	Year Ended December 31,	
	2013	2012
Net cash used in operating activities	\$ (9,949)	\$ (9,730)
Net cash provided by investing activities	253	4,275
Net cash provided by financing activities	9,234	4,865
Net cash used	\$ (462)	\$ (590)

Net Cash Used in Operating Activities

Net cash used in operating activities was \$9.9 million in 2013, compared to \$9.7 million used in 2012, an increase of \$0.2 million, or 2%. The cash used in operating activities for 2013 was primarily attributable to a \$4.2 million increase in the change in prepaid expenses and other current assets primarily due to the deferred offering costs and a \$2.6 million increase in the change in inventory as we built up our inventory. These amounts were partially offset by a \$1.2 million decrease in trade accounts receivable mostly due to improved collection and cash management efforts and a \$3.3 million increase in accounts payable and accrued liabilities.

Net Cash Provided by Investing Activities

Net cash provided by investing activities was \$0.3 million in 2013, compared to \$4.3 million provided in 2012, a decrease of \$4.0 million, or 93%. The decrease in net cash provided by investing activities was primarily attributable to a \$7.5 million decrease in the proceeds from maturities of marketable securities and a \$1.7 million increase in the purchase of property and equipment, which were partially offset by a \$5.1 million decrease in purchases of marketable securities.

Net Cash Provided by Financing Activities

Net cash provided by financing activities was \$9.2 million in 2013, compared to \$4.9 million provided in 2012, an increase of \$4.3 million, or 90%. This increase in net cash provided by financing activities was primarily attributable to an \$8.9 million increase in net proceeds from the issuance of convertible preferred stock and a \$2.9 million increase in proceeds from the issuance of common stock in connection with the exercise of common stock warrants and options, partially offset by a \$4.8 million decrease in net proceeds from the issuance of convertible debt and a \$3.1 million increase in net payments on our GE Secured Lending Facility.

Indebtedness

In December 2012, we entered into the GE Secured Lending Facility, which consists of a \$18.0 million term loan and up to \$3.5 million revolving credit facility with GE Capital, as agent and lender, and Zions First National Bank, as lender. We pledged all of our assets as collateral for the loans. The revolving line of credit is secured by our accounts receivable, based on certain defined criteria. The term loan consisted of interest only payments until January 1, 2014. Beginning in January 2014, monthly interest payments as well as principal payments of approximately \$600,000 each are required for a period of 30 months. We were in default of the liquidity covenant under the GE Secured Lending Facility in November 2013, and, in December 2013, we amended the terms of the GE Secured Lending Facility to allow for a temporary waiver effective from November 1, 2013 through January 31, 2014 of the liquidity

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covenant under the agreement discussed below. In addition, we agreed to increase the credit reserve from \$0.5 million to \$1.0 million. On January 28, 2014, we obtained an additional waiver of the liquidity covenant from GE Capital through February 28, 2014 and agreed to increase the credit reserve under this facility by an additional \$0.5 million, bringing the total reserve to \$1.5 million. The total accrued amendment fees at December 31, 2013 were \$1.1 million. We also agreed to pay GE Capital a fee of \$200,000 in connection with the January 28, 2014 waiver if the GE Secured Lending Facility is not repaid on or before March 31, 2014. We expect to pay the \$1.3 million in amendment fees in April 2014.

The term loan bears interest at the fixed rate of 7.5% per annum, while the line of credit had an interest rate of 7.0% at December 31, 2013, which is based on the variable rate of 5.5% plus the higher of (i) 1.5% and (ii) the three-month LIBOR, determined as of two London business days divided by a number equal to 1.0 minus the aggregate of the rates of reserve requirements on the day that is two London business days prior to the beginning of the interest period for Eurocurrency funding that are required to be maintained by a member bank of the Federal Reserve System. The agreement includes a non-refundable final payment fee equal to 4% of the original principal amount of the term loan, or \$720,000, upon prepayment in full or scheduled maturity of the term loan, as well as an annual management fee equal to \$15,000 per year.

The loan agreement includes certain financial covenants related to monthly cash burn and minimum liquidity, days sales outstanding of accounts receivable balances, annual payment restrictions to our directors and other financial reporting requirements. The liquidity covenant requires us to maintain cash and cash equivalents and availability under the revolving credit facility equal to the greater of \$1.5 million (exclusive of availability under the revolving credit facility) or six times our monthly cash burn, as defined in the revolving credit facility. This covenant may significantly limit our ability to use our cash and cash equivalents to fund our operations. The loan agreement provides for an unused credit facility fee of 0.75% per annum of the unused portion of the line of credit, payable monthly in arrears.

We paid a total of approximately \$333,000 in fees and commissions associated with entering into this facility, of which approximately \$264,000 was capitalized as debt issuance costs and the remaining \$69,000 was recorded as interest expense in 2012.

In connection with entering into the GE Secured Lending Facility, we repaid all amounts outstanding under our term loans and line of credit facility with a previous lender in December 2012, which totaled \$18.0 million in principal and approximately \$36,000 in accrued interest. We paid \$107,500 in commissions related to this repayment, of which approximately \$70,000 was capitalized as debt issuance costs and the remaining \$37,500 was recorded as interest expense in 2012. We used a portion of the net proceeds of the IPO to service the outstanding debt under the facility.

Contractual Obligations and Commitments

The following table summarizes our outstanding contractual obligations as of December 31, 2013 (in thousands):

	Total	Less Than 1 Year	1-3 Years	4-5 Years	After 5 Years
Long-term debt (1)	\$18,000	\$ 7,200	\$10,800	\$ —	\$ —
Operating leases	5,529	856	1,793	1,900	980
Total contractual obligations	\$23,529	\$ 8,056	\$12,593	\$ 1,900	\$ 980

(1) Does not include the \$720,000 final payment fee we must pay upon prepayment in full or scheduled maturity of the term loan, the \$15,000 per year annual management fee, the amendment fees of up to \$1.1 million, which are due at the earlier of the date the debt is paid in full or March 31, 2014, or monthly interest payments.

The information above reflects only payment obligations that are fixed and determinable. Our commitments for long-term debt relate to our term loans with GE Capital and our commitment to our operating lease for our corporate headquarters and manufacturing facility in Salt Lake City, Utah. The above table does not include any of the contractual obligations with respect to royalties payable upon sales of certain of our products as none of our arrangements contain minimum royalty payments. We also do not have contractually minimum purchase commitments for the supply of any of our raw materials, products or instruments.

Off-Balance Sheet Arrangements

We do not have any off-balance sheet arrangements, as defined in Item 303(a)(4) of Regulation S-K.

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Related-Party Transactions

For a description of our related-party transactions, see “Certain Relationships and Related Party Transactions.”

Seasonality and Backlog

Our business is generally not seasonal in nature. However, our sales may be influenced by summer vacation and winter holiday periods during which we believe fewer spinal fusion surgeries are conducted. Our sales generally consist of products that are in stock with us or maintained at hospitals or with our sales distributors. Accordingly, we do not have a backlog of sales orders.

Critical Accounting Policies and Estimates

The preparation of the consolidated financial statements requires us to make assumptions, estimates and judgments that affect the reported amounts of assets and liabilities, the disclosures of contingent assets and liabilities as of the date of the consolidated financial statements, and the reported amounts of product revenues and expenses during the reporting periods. Certain of our more critical accounting policies require the application of significant judgment by management in selecting the appropriate assumptions for calculating financial estimates. By their nature, these judgments are subject to an inherent degree of uncertainty. On an ongoing basis, we evaluate our judgments, including those related to inventories, recoverability of long-lived assets and the fair value of our common stock. We use historical experience and other assumptions as the basis for our judgments and making these estimates. Because future events and their effects cannot be determined with precision, actual results could differ significantly from these estimates. Any changes in those estimates will be reflected in our consolidated financial statements as they occur. As an “emerging growth company,” we have elected to delay the adoption of new or revised accounting standards until those standards would otherwise apply to private companies. As a result, our financial statements may not be comparable to those of other public companies. While our significant accounting policies are more fully described in the footnotes to our consolidated financial statements included elsewhere in this Annual Report, we believe that the following accounting policies and estimates are most critical to a full understanding and evaluation of our reported financial results. The critical accounting policies addressed below reflect our most significant judgments and estimates used in the preparation of our consolidated financial statements.

Revenue Recognition

We derive our product revenue primarily from the sale of spinal fusion devices and related products used in the treatment of spine disorders. Our product revenue is generated from sales to two types of customers: (1) surgeons and hospitals; and (2) stocking distributors. Most of our products are sold on a consignment basis through a network of independent sales distributors; however, we also sell our products to independent stocking distributors. Product revenue is recognized when all four of the following criteria are met: (1) persuasive evidence that an arrangement exists; (2) delivery of the products has occurred; (3) the selling price of the product is fixed or determinable; and (4) collectability is reasonably assured. We generate the majority of our revenue from the sale of inventory that is consigned to independent sales distributors that sell our products to surgeons and hospitals. For these products, we recognize revenue at the time we are notified the product has been used or implanted and a valid purchase order has been received. For all other transactions, we recognize revenue when title and risk of loss transfer to the stocking distributor, and all other revenue recognition criteria have been met. We generally recognize revenue from sales to stocking distributors at the time the product is shipped to the distributor. Stocking distributors, who sell the products to their customers, take title to the products and assume all risks of ownership at time of shipment. Our stocking distributors are obligated to pay within specified terms regardless of when, if ever, they sell the products. Our policy is to classify shipping and handling costs billed to customers as an offset to total shipping expense in the statement of operations, primarily within sales and marketing. In general, our customers do not have any rights of return or exchange.

Accounts Receivable and Allowance for Doubtful Accounts

The majority of our accounts receivable is composed of amounts due from hospitals or surgical centers. Accounts receivable are carried at cost less an allowance for doubtful accounts. On a regular basis, we evaluate accounts receivable and estimate an allowance for doubtful accounts, as needed, based on various factors such as customers’ current credit conditions, length of time past due, and the general economy as a whole. Receivables are written off against the allowance when they are deemed uncollectible.

Inventories

Inventories are stated at the lower of cost or market, with cost for manufactured inventory determined under the standard cost method which approximates the first-in first-out method. Manufactured inventory consists of raw material, direct labor and manufacturing overhead cost components. Inventories purchased from third-party manufacturers are stated at the lower of cost or market using the first-in, first out method. We review the carrying value of inventory on a periodic basis for excess or obsolete items

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and record an expense for the identified items as necessary. We have made adjustments to, and it is reasonably possible that we may be required to make further adjustments to, the carrying value of inventory in future periods. We hold some consigned inventory at distributors and other customer locations where revenue recognition criteria have not yet been met.

Long-Lived Assets, Indefinite-Lived Intangibles and Goodwill

Periodically we assess potential impairment of our long-lived assets, which include property, equipment, and acquired intangible assets. We perform an impairment review whenever events or changes in circumstances indicate that the carrying value may not be recoverable. Factors we consider important which could trigger an impairment review include, but are not limited to, significant under-performance relative to historical or projected future operating results, significant changes in the manner of use of the acquired assets or our overall business strategy, and significant industry or economic trends. When we determine that the carrying value of a long-lived asset may not be recoverable based upon the existence of one or more of the above indicators, we determine the recoverability by comparing the carrying amount of the asset to net future undiscounted cash flows that the asset is expected to generate and recognize an impairment charge equal to the amount by which the carrying amount exceeds the fair market value of the asset. We amortize intangible assets on a straight-line basis over their estimated useful lives.

For indefinite lived intangible assets that are not subject to amortization, the impairment test consists of a comparison of the fair value of an intangible asset with its carrying amount. If the carrying amount of an intangible asset exceeds its fair value, an impairment loss is recognized in an amount equal to that excess.

Our management noted that certain US Spine product sales and sales to certain acquired US Spine customers during the one-year period ended December 31, 2012 had been less than expected relative to the forecasted revenues at the time of our acquisition of US Spine. This indicator prompted us to question whether the carrying value of our long-lived and indefinite lived intangible assets would be recoverable. We compared the carrying amount of the assets to net future undiscounted cash flows that the intangible assets are expected to generate, and concluded that an impairment existed. We estimated the fair values of the intangible assets and recognized an impairment loss of approximately \$15.3 million in the year ended December 31, 2012. No impairment loss was recognized in the year ended December 31, 2013. Should conditions change such that our estimates of associated undiscounted cash flows would not support the unamortized carrying value of specific assets, we could have further impairments.

The income approach used in our 2012 impairment analysis considered management's business plans and projections as the basis for expected cash flows for the next ten years and a 5% residual growth rate thereafter. We also used a weighted average discount rate of 17%, a weighted average revenue growth rate ranging from (58)% to 10% and an EBITDA margin ranging from approximately 9% to 12.4% for the analysis.

Our long-lived assets include surgical instruments used by spine surgeons during surgical procedures to facilitate the implantation of our products. There are no contractual terms with respect to the usage of our instruments by our customers. Surgeons are under no contractual commitment to use our instruments. We maintain ownership of these instruments and, when requested, we allow the surgeons to use the instruments to facilitate implantation of our related products. We do not currently charge for the use of our instruments and there are no minimum purchase commitments of our products. As our surgical instrumentation is used numerous times over several years, often by many different customers, instruments are capitalized as property and equipment once they have been placed in service. Once placed in service, instruments are carried at cost, less accumulated depreciation. Depreciation is computed using the straight-line method based on average estimated useful lives. Estimated useful lives of surgical instruments are three years and are determined based on a variety of factors including reference to associated product life cycles. As instruments are used as tools to assist surgeons, depreciation of instruments is recognized as a sales and marketing expense. Instrument depreciation expense was \$1.0 million and \$1.2 million for the years ended December 31, 2013 and 2012, respectively.

We review our long-lived assets for impairment whenever events or changes in circumstances indicate that the carrying value of the assets may not be recoverable. An impairment loss would be recognized when estimated future undiscounted cash flows relating to the assets are less than the assets' carrying amount. An impairment loss is measured as the amount by which the carrying amount of an asset exceeds its fair value.

We test goodwill for impairment annually as of December 31, or whenever events or changes in circumstances indicate that goodwill may be impaired. We initially assess qualitative factors to determine whether the existence of events or circumstances leads to a determination that it is more-likely-than-not that the fair value of a reporting unit is less than its carrying amount. For goodwill impairment testing purposes, we consider the value of our equity, including the value of our convertible preferred stock, in the total carrying value of our single reporting unit. If, after assessing the totality of events or circumstances, we determine it is more-likely-than-not that the fair value of our reporting unit is less than its carrying amount, then we perform a first step analysis by comparing the carrying amount of net assets to the fair value of our single reporting unit. If the fair value is determined to be less than the carrying amount, a second step analysis is performed to compute the amount of impairment as the difference between the implied estimated fair value of goodwill and the carrying amount.

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At December 31, 2013, the balance of goodwill resulting from the US Spine acquisition was \$6.2 million. We measure the fair value of our reporting unit for purposes of our impairment test utilizing the income approach. The income approach is calculated based on management's best estimates of future cash flows which depend primarily upon revenue growth, discount rate, terminal value and long-term growth rate and total operating expenses. There is a certain degree of uncertainty associated with these key assumptions and there are potential events and circumstances that could reasonably be expected to affect these key assumptions, such as (i) significant decline in product revenue or failure to increase revenue in future years, (ii) failure of the new Design and Build Program to increase revenue as expected, (iii) significant increases in the manufacturing costs or acquisition costs of our inventory and (iv) lack of clearance or approval from the FDA for any of our future product candidates.

Income Taxes

We recognize deferred tax assets and liabilities for the future tax consequences attributable to the differences between the financial statement carrying value of existing assets and liabilities and their respective tax bases. Deferred tax assets and liabilities are measured using enacted tax rates in effect for the fiscal year in which those temporary differences are expected to be recovered or settled. Valuation allowances are established when necessary to reduce deferred tax assets to the amount expected to be realized.

We operate in various tax jurisdictions and are subject to audit by various tax authorities. We provide for tax contingencies whenever it is deemed probable that a tax asset has been impaired or a tax liability has been incurred for events such as tax claims or changes in tax laws. Tax contingencies are based upon their technical merits relative tax law and the specific facts and circumstances as of each reporting period. Changes in facts and circumstances could result in material changes to the amounts recorded for such tax contingencies.

We recognize uncertain income tax positions taken on income tax returns at the largest amount that is more-likely than-not to be sustained upon audit by the relevant taxing authority. An uncertain income tax position will not be recognized if it has less than a 50% likelihood of being sustained.

Our policy for recording interest and penalties associated with uncertain tax positions is to record such items as a component of our income tax provision. For the years ended December 31, 2013 and 2012, we did not record any material interest income, interest expense or penalties related to uncertain tax positions or the settlement of audits for prior periods.

Stock-Based Compensation Expense

Common Stock Valuation

Historically, our board of directors has determined the fair value of the common stock with assistance from management and based upon information available at the time of grant. The valuation of our common stock requires us to make complex and subjective judgments. We considered a combination of valuation methodologies, including income, market and transaction approaches. The most significant factors considered by our board of directors when determining the fair value of our common stock were as follows:

- external market and economic conditions affecting the medical device industry;
- prices at which we sold shares of our convertible preferred stock to third-party investors;
- the superior rights and preferences of securities senior to our common stock, such as our preferred stock, at the time of each grant;
- our need for future financing to fund commercial operations;
- the lack of marketability of our common stock;
- third-party valuations of our common stock;
- our historical operating and financial performance;
- the status of our research and development efforts;
- the status of our new product releases to the spine market;
- the likelihood of achieving a liquidity event, such as an initial public offering or sale of our company; and
- estimates and analysis provided by management.

We have regularly obtained third-party valuations to assist our board of directors in determining the fair value of our common stock for each stock option grant and other stock-based awards, on an annual basis since 2007.

Significant Factors and Assumptions Used in Determining Fair Value of Common Stock

The \$5.75 issuance price of the common stock sold in our IPO in February 2014 was used as the estimated fair value of the shares of our common stock as of and for the quarter ended December 31, 2013 due to the close proximity to the valuation dates and there

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were no significant changes that would indicate a decrease or increase in the stock price. The price of \$17.53 was used as the estimated fair value of our common shares for estimated fair value calculations done as of and prior to September 30, 2013. The price of \$17.53 was based on a third party valuation performed in September 2012. The valuation technique used in the third party valuation was a hybrid of the discounted cash flow method and the guideline public company method. The significant assumptions used in determining the \$17.53 estimated fair value of our common shares using a hybrid of the discounted cash flow method and the guideline public company methodology were as follows:

Weighted-average cost of capital (WACC)	17%
Revenue growth rate (range)	32.5% to 5%
Compounded average revenue growth rate	17.7%
EBITDA margin (range)	(23.8)% to 32.7%

Stock-Based Compensation

We apply the fair value recognition provisions of Financial Accounting Standards Board, or FASB, Accounting Standards Codification, or ASC, Topic 718, *Compensation-Stock Compensation*, or ASC 718. Determining the amount of stock-based compensation to be recorded requires us to develop estimates of the fair value of stock options and other equity awards as of their grant date. Stock-based compensation expense is recognized ratably over the requisite service period, which in most cases is the vesting period of the award. Calculating the fair value of stock-based awards requires that we make highly subjective assumptions. Use of this valuation methodology requires that we make assumptions as to the volatility of our common stock, the expected term of our stock options, the risk free rate of return for a period that approximates the expected term of our stock options and our expected dividend yield. Because we are a privately-held company with no trading history, we utilize the historical stock price volatility from a representative group of public companies to estimate expected stock price volatility. We selected companies from the medical device industry, specifically those who are focused on the design, development and commercialization of products for the treatment of spine disorders, and who have similar characteristics to us, such as stage of life cycle and size. We intend to continue to utilize the historical volatility of the same or similar public companies to estimate expected volatility until a sufficient amount of historical information regarding the price of our publically traded stock becomes available. We use the simplified method as prescribed by the Securities and Exchange Commission Staff Accounting Bulletin No. 107, *Share-based Payment*, to calculate the expected term of stock option grants to employees as we do not have sufficient historical exercise data to provide a reasonable basis upon which to estimate the expected term of stock options granted to employees. We utilize a dividend yield of zero because we have never paid cash dividends and have no current intention to pay cash dividends. The risk-free rate of return used for each grant is based on the U.S. Treasury yield curve in effect at the time of grant for instruments with a similar expected life. No options were granted to employees during the year ended December 31, 2013. The following assumptions were used in the calculation to estimate the fair value of options granted to employees using the Black-Scholes-Merton valuation model during the year ended December 31, 2012:

Weighted-average risk-free interest rate	1.14%
Weighted-average expected life (in years)	5.34
Expected dividend yield	0%
Weighted-average expected volatility	72%

No options were granted to non-employees during the year ended December 31, 2012. The following assumptions were used in the Black-Scholes-Merton valuation model related to non-employee stock options granted during the year ended December 31, 2013:

Weighted-average risk-free interest rate	2.74%
Weighted-average expected life (in years)	10.0
Expected dividend yield	0%
Weighted-average expected volatility	52%

The estimated fair value of stock-based awards for employee and non-employee director services are expensed over the requisite service period. Option awards issued to non-employees, excluding non-employee directors, are recorded at their fair value as determined in accordance with authoritative guidance, are periodically revalued as the options vest and are recognized as expense over the related service period. As a result, the charge to operations for non-employee awards with vesting conditions is affected each reporting period by changes in the fair value of our common stock.

We are required to estimate the level of forfeitures expected to occur and record stock-based compensation expense only for those awards that we ultimately expect will vest. We estimate our forfeiture rate based on the type of award, employee class and historical experience. Through December 31, 2013, actual forfeitures have not been material.

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In February 2013, our employees elected to exchange 93,968 options to purchase our common stock for restricted stock units, or RSUs, pursuant to a one-time tender offer authorized by our board of directors. The RSUs were issued under the 2012 Plan and have three-year terms and vest upon the earlier of a change in control or upon the expiration of the lock-up period for our IPO on August 12, 2014. The incremental fair value on the date of the exchange between the original stock options and the RSUs issued of approximately \$0.8 million will be recognized over a period of six months commencing in February 2014.

At December 31, 2013, we had 168,832 unvested RSUs, the majority of which will vest upon the earlier of a change in control or on August 12, 2014, the expiration of the lock-up period for our IPO. Subsequent to December 31, 2013, our board approved grants on January 27, 2014 totaling 1,692,980 RSUs to be issued on effectiveness of the filing of a registration statement on Form S-8 and vest upon the expiration of the lock-up period for our IPO. We will take compensation charges upon vesting of RSUs based upon their grant date fair value. The estimated aggregate value to be recognized as compensation expense in 2014 for the RSUs granted on January 27, 2014 and outstanding RSUs at December 31, 2013 is expected to be approximately \$11.4 million, including the \$0.8 million expense as a result of the exchange described above.

Total stock-based compensation expense totaled \$0.6 million and \$1.0 million for 2013 and 2012, respectively. As of December 31, 2013, we had approximately \$2.0 million of total unrecognized stock-based compensation expense. While our stock-based compensation to date has not been material to our financial results, we expect the impact to grow in future periods due to the issuance of RSUs in 2013 and 2014 for which minimal expense has been recorded to date, and the potential increases in the value of our common stock and headcount.

Common Stock Warrant Liability and Preferred Stock Warrant Liability

As of December 31, 2013, we had warrants outstanding to purchase shares of our Series C, Series D, Series E and Series F convertible preferred stock and common stock. Freestanding warrants that are related to the purchase of redeemable preferred stock are classified as liabilities and recorded at fair value regardless of the timing of the redemption feature or the redemption price or the likelihood of redemption. The warrants are subject to re-measurement at each balance sheet date and any change in fair value is recognized as a component of other income (expense), net in our statement of comprehensive loss. We measure the fair value of our warrants to purchase our convertible preferred stock using a Black-Scholes-Merton option pricing model. The warrants to purchase shares of our common stock contain a provision requiring a reduction to the exercise price in the event we issue common stock, or securities convertible into or exercisable for common stock, at a price per share lower than the warrant exercise price. The anti-dilution feature requires the warrants to be classified as liabilities and re-measured at fair value at each balance sheet date. The fair value of the warrants to purchase common stock on the date of issuance and on each re-measurement date is classified as a liability and is estimated using the Black-Scholes-Merton valuation model. Any modifications to the warrant liabilities are recorded in earnings during the period of the modification. The significant assumptions used in estimating the fair value of our warrant liabilities include the exercise price, volatility of the stock underlying the warrant, risk-free interest rate, estimated fair value of the stock underlying the warrant and the estimated life of the warrant.

Upon consummation of our IPO in February 2014, all classes of our convertible preferred stock were converted into common stock. Upon conversion of the underlying classes of convertible preferred stock, pursuant to the terms of the preferred stock warrants, the remaining warrants to purchase our Series C, Series D, Series E and Series F convertible preferred stock were classified as a component of equity and no longer be subject to re-measurement. However, the common stock warrant liability will continue to be required to be re-measured at each balance sheet date, until such time that the common stock warrants are exercised or expire.

Recently Issued Accounting Pronouncements

In February 2013, the FASB issued an update to improve the transparency of reporting reclassifications out of accumulated other comprehensive income. The amendments in the update did not change the current requirements for reporting net income or other comprehensive income in financial statements. The new amendments require an organization to present (either on the face of the statement where net income is presented or in the notes) the effects on the line items of net income of significant amounts reclassified out of accumulated other comprehensive income if the item reclassified is required under generally accepted accounting principles in the United States, or U.S. GAAP, to be reclassified to net income in its entirety in the same reporting period. Additionally, for other amounts that are not required under U.S. GAAP to be reclassified in their entirety to net income in the same reporting period, an entity is required to cross-reference other disclosures required under U.S. GAAP to provide additional detail about those amounts. The amendments were effective for the year ended December 31, 2013. We have adopted and the new guidance did not have a material impact on the consolidated financial statements.

Jumpstart Our Business Startups Act of 2012

On April 5, 2012, the Jumpstart Our Business Startups Act of 2012, or JOBS Act, was enacted. Section 107 of the JOBS Act, provides that an “emerging growth company” can take advantage of the extended transition period provided in Section 7(a)(2)(B) of

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the Securities Act of 1933, as amended, for complying with new or revised accounting standards. In other words, an “emerging growth company” can delay the adoption of certain accounting standards until those standards would otherwise apply to private companies. We are electing to delay such adoption of new or revised accounting standards, and as a result, we may not comply with new or revised accounting standards on the relevant dates on which adoption of such standards is required for non-emerging growth companies. As a result of this election, our financial statements may not be comparable to the financial statements of other public companies. We may take advantage of these reporting exemptions until we are no longer an “emerging growth company.”

We are in the process of evaluating the benefits of relying on other exemptions and reduced reporting requirements provided by the JOBS Act. Subject to certain conditions set forth in the JOBS Act, as an “emerging growth company,” we intend to rely on certain of these exemptions, including without limitation, (1) providing an auditor’s attestation report on our system of internal controls over financial reporting pursuant to Section 404(b) of the Sarbanes-Oxley Act and (2) complying with any requirement that may be adopted by the Public Company Accounting Oversight Board regarding mandatory audit firm rotation or a supplement to the auditor’s report providing additional information about the audit and the consolidated financial statements, known as the auditor discussion and analysis. We may be able to remain an “emerging growth company” until the earliest of (a) the last day of the fiscal year in which we have total annual gross revenues of \$1 billion or more, (b) the last day of our fiscal year following the fifth anniversary of the date of our IPO, (c) the date on which we have issued more than \$1 billion in non-convertible debt during the previous three years or (d) the date on which we are deemed to be a large accelerated filer under the rules of the SEC. Additionally, we are also currently a “smaller reporting company” as defined in the Securities Exchange Act of 1934, and in the event that we are still considered a smaller reporting company at such time as we cease being an emerging growth company, we will be exempt from the provisions of Section 404(b) of the Sarbanes-Oxley Act requiring that independent registered public accounting firms provide an attestation report on the effectiveness of internal control over financial reporting.

ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

Not applicable.

ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA

Financial Statements

The consolidated financial statements of Amedica appear at the end of this Annual Report beginning with the index to Financial Statements on page F-1 (see Part IV, Item 15 “Financial Statements”), and are incorporated herein.

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

Not applicable.

ITEM 9A. CONTROLS AND PROCEDURES

(a) Disclosure Controls and Procedures

We maintain a set of disclosure controls and procedures (as defined in Rule 13a-15(e) of the Exchange Act) designed to ensure that information required to be disclosed in reports filed or submitted under the Exchange Act is recorded, processed, summarized, and reported within the time periods specified in rules and forms adopted by the SEC.

In accordance with Rule 13a-15(b) of the Exchange Act, as of the end of the period covered by this Annual Report on Form 10-K, an evaluation was carried out under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, to assess the effectiveness of our disclosure controls and procedures, as of the end of the period covered by this Annual Report on Form 10-K were effective to provide reasonable assurance that information required to be disclosed by us in reports that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the SEC’s rules and forms and is accumulated and communicated to our management, including the Chief Executive Officer and Chief Financial Officer, as appropriate to allow timely decisions regarding required disclosure.

(b) Management’s Report on Internal Control over Financial Reporting and Attestation Report of Registered Public Accounting Firm

This Annual Report does not include a report of management’s assessment regarding internal control over financial reporting or an attestation report of our registered public accounting firm due to a transition period established by rules of the SEC applicable to emerging growth companies.

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(c) Changes in Internal Control Over Financial Reporting

There were no changes in our internal control over financial reporting that occurred during the quarter ended December 31, 2013 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

ITEM 9B. OTHER INFORMATION

None.

PART III

ITEM 10. DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE

Directors and Executive Officers

Our current directors and executive officers and their respective ages and positions are as follows:

<u>Name</u>	<u>Age</u>	<u>Position</u>
Max E. Link, Ph.D.	73	Chairman of the Board of Directors
Eric K. Olson	50	Director, President and Chief Executive Officer
Jay M. Moyes	60	Director and Chief Financial Officer
B. Sonny Bal, M.D.	51	Director
David W. Truetzel	57	Director
Jeffrey S. White	60	Director
James P. Abraham	54	Senior Vice President, Global Sales
Kevin Davis	48	Chief Operating Officer
Bryan J. McEntire	62	Chief Technology Officer
Kevin Ontiveros	53	Chief Legal Officer, Chief Compliance Officer and Corporate Secretary
Vytas Rupinkas	61	Vice President, Marketing
Christopher R. Whitfield	46	Chief Commercial Officer

The following is a brief summary of the background of each of our current directors and executive officers.

Max E. Link, Ph.D. has served as the chairman of our board of directors since October 2003. Dr. Link was chairman of the board of directors and Chief Executive Officer of Centerpulse AG, a medical implant company from March 2002 to October 2003. Prior to joining Centerpulse, Dr. Link was Chief Executive Officer of Corange (Bermuda), the parent company of Boehringer Mannheim Corporation and chairman of the board of directors and chief executive officer of Sandoz Pharma, Ltd., now part of Novartis Corporation, a manufacturer of pharmaceutical products. Dr. Link is chairman of the boards of directors of three publicly listed biopharmaceutical companies, Alexion Pharmaceuticals, Inc., Celsion Corporation and CytRx Corporation. Dr. Link holds a Ph.D. in Economics from University of St. Gallen (Switzerland).

We believe that Dr. Link is qualified to serve as a member of our board of directors because of his significant experience leading companies in our industry as well as the depth of his institutional knowledge of our company.

Eric K. Olson has served as our Chief Executive Officer and President and as a director since February 2012. Prior to serving us in this capacity, Mr. Olson served as our Senior Vice President of Global Marketing from June 2011 through February 2012. From December 2007 to June 2011, Mr. Olson was the Executive Vice President of Sales & Marketing for Axial Biotech, Inc., a molecular diagnostics company. Mr. Olson has also held senior sales and marketing positions with Medtronic, Inc. and Smith & Nephew. Mr. Olson holds a B.S. in Behavioral Science and Health Administration from the University of Utah, and has also completed a master's-level internship program at the same institution.

We believe that Mr. Olson's position as the Chief Executive Officer and President of our Company uniquely qualifies him to serve on our board of directors due to his intimate knowledge of our day-to-day operations. Additionally, Mr. Olson possesses a wealth of industry experience related to our business.

Jay M. Moyes has served on our board of directors since November 2012 and as our Chief Financial Officer since October 2013. Since November 2007 Mr. Moyes has been the managing member of Drayton Investments, LLC, a partnership focused on investing in private healthcare related companies and real estate financing. In April 2012, he joined the board of directors of Puma Biotechnology Inc., a biopharmaceutical company. Since May 2006, he has been a member of the board of directors and chairman of the audit committee of Osiris Therapeutics, Inc., a publicly held stem cell therapeutics company. Mr. Moyes is also a director of BioCardia, Inc., a medical device company, and Integrated Diagnostics Inc., a molecular diagnostics company. From May 2008 through July 2009, Mr. Moyes served as the Chief Financial Officer of XDx, Inc., a privately held molecular diagnostics company. Prior to that, he served as the Chief Financial Officer of Myriad Genetics, Inc., a publicly held healthcare diagnostics company, from June 1996 until his retirement in November 2007, and as its Vice President of Finance from July 1993 until July 2005. From 1991 to 1993, Mr. Moyes served as Vice President of Finance and Chief Financial Officer of Genmark, Inc., a privately held genetics company. He held various positions with the accounting firm of KPMG LLP from 1979 through 1991, most recently as a Senior Manager. He also served as a member of the Board of Trustees of the Utah Life Science Association from 1999 through 2006. He holds an M.B.A from the University of Utah and received his B.A. in Economics from Weber State University.

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In addition to serving as our Chief Financial Officer, we believe that Mr. Moyes' experience working with biotechnology companies through their transformation from emerging growth to established, publicly-traded companies qualify him to serve on our board of directors.

B. Sonny Bal, M.D. has served on our board of directors since February 2012. Dr. Bal is Professor & Chief of Adult Reconstruction at the University of Missouri, Columbia, specializing in hip and knee replacement surgery. He also is an Adjunct Professor of Material Sciences at the University of Missouri at Rolla. Dr. Bal is a member of the American Academy of Orthopaedic Surgeons and the American Association of Hip and Knee Surgeons. Dr. Bal received his M.D. degree from Cornell University and an M.B.A. from Northwestern University, and a J.D. from the University of Missouri. Dr. Bal is a licensed attorney and co-founder of the Bal Brenner law firm in North Carolina.

We believe that Dr. Bal's expertise in orthopedic surgery and his specialty in hip and knee replacement surgery qualifies him to serve on our board of directors.

David W. Truetzel has served on our board of directors since our acquisition of US Spine, Inc. in September 2010. Mr. Truetzel has been the general partner of Augury Capital Partners a private equity fund that invests in life sciences and information technology companies, which he co-founded in 2006. Mr. Truetzel is a director of Enterprise Bank, Inc., Verifi, Inc., a provider of electronic payment solutions, Clearent, LLC, a credit card processing provider, and Paranet, LLC, an IT services provider. Mr. Truetzel holds a B.S. in Business Administration from Saint Louis University, an M.B.A. from The Wharton School and is a licensed C.P.A.

We believe that Mr. Truetzel's financial and managerial expertise qualify him to serve on our board of directors.

Jeffrey S. White has served on our board of directors since January 2014. Since January 2013, Mr. White has served as Principal at Medtech Advisory Group LLC, a firm he founded that advises early and mid-stage medical technology firms. Mr. White is currently a director of Residency Select LLC, a company which offers psychometric assessment, training and compliance products to medical and surgical residency programs. From May 2006 to December 2012 he served as Global Director of Business Development for Synthes Inc., a global orthopedic firm that was acquired by Johnson and Johnson in 2012. Mr. White has served as Chief Executive Officer and co-founder of several start-up surgical device firms and has previously held executive level positions at Richard-Allan Medical Industries Inc., a medical device manufacturer, which was acquired by Urohealth Systems Inc. and United States Surgical Corporation, unit of Covidien plc. Mr. White holds a B.S. in Biology from Union College in Schenectady NY.

We believe that Mr. White's experience as an executive and founder of medical device companies qualifies him to serve on our board of directors.

James P. Abraham joined us as Senior Vice President, Global Sales, in January 2013. From January 2007 to December 2013 Mr. Abraham worked in various capacities at Stryker Corporation, a medical equipment company including as Senior Director of Sales. He also previously served as Senior Vice President of Sales and Marketing for IsoTis Orthobiologics, Inc., a company which specializes in human tissue and synthetic grafting and injectable bone growth stimulation. Mr. Abraham holds a B.S. in Business Administration from Creighton University.

Kevin Davis has served as our Chief Operating Officer since June 2012. From December 2011 to June 2012, he served as our President of Manufacturing. From March 2011 to December 2011, he served as our Vice President of Strategy and Business Development. From March 2009 to March 2011, he served as our Cost Accountant, Financial Systems. Mr. Davis was the Chief Financial Officer, from April 2007 to March 2009, of Nevada Chemicals, Inc., a sodium cyanide chemical company and served as one of its directors. Mr. Davis graduated from the University of Utah with a B.S. in Accounting.

Bryan J. McEntire has served as our Chief Technology Officer since May 2012. From June 2004 to May 2012 he served as our Vice President of Manufacturing and as our Vice President of Research from December 2006 to May 2012. Mr. McEntire has worked in various advanced ceramic product development, quality engineering and manufacturing roles at Applied Materials, Inc., Norton Advanced Ceramics, a division of Saint-Gobain Industrial Ceramics Corporation, Norton/TRW Ceramics and Ceramatec, Inc., a small producer of ionic-conducting and structural ceramic components located in Salt Lake City, Utah. Mr. McEntire holds a B.S. degree in Materials Science and Engineering and an M.B.A. from the University of Utah.

Kevin Ontiveros has served as our Chief Legal Officer and Chief Compliance Officer since December 2012. Mr. Ontiveros was previously a practicing attorney at Life Science Law PC from February 2011 to December 2012 and Stoel Rives LLP from January 2009 to January 2011, where he provided legal and business counsel on a wide range of matters, including technology licensing transactions, corporate financing opportunities (including public and private equity and debt offerings), public company SEC reporting

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compliance, and clinical trial, manufacturing, distribution, and research and development agreements. Mr. Ontiveros served as the Vice President-Legal Affairs, General Counsel and Corporate Secretary for ImaRx Therapeutics, Inc. from March 2007 to December 2008 and as the Vice President-Corporate Law and Assistant Corporate Secretary for NPS Pharmaceuticals, Inc. from April 1996 to March 2007. Mr. Ontiveros earned his B.A. from the University of Arizona, his J.D. from the University of Utah School of Law and his L.L.M. in Taxation from the University of Florida.

Vytas Rupinskas is our Vice President of Marketing, a position he has held since December 2012. From September 2005 to March 2012, Mr. Rupinskas served as the Director of Product Management for Leads & Accessories for the Neuromodulation Division of St. Jude Medical, Inc., and Marketing Manager for St. Jude Medical, a provider of implantable medical devices. Prior to his tenure at St. Jude Medical, Mr. Rupinskas served as Senior Product Manager at Exactech, Inc., a provider of implant devices and surgical instruments and held various senior global marketing and international sales management positions at DePuy Orthopaedics, Inc., a medical device company, and its affiliates DePuy International Ltd. and DePuy Spine, Inc. Mr. Rupinskas is a graduate of the University of Illinois with a B.S. degree in Liberal Arts and Sciences and an M.S. in Mechanical Engineering.

Christopher R. Whitfield has served as our Chief Commercial Officer since November 2013. From March 2012 to September 2012, Mr. Whitfield served as the Executive Vice President, Sales and Marketing of Pioneer Surgical Technologies and from October 2009 to March 2012 as its Vice President, Marketing. From October 2008 to September 2009, he served as the West Area Vice President, Sales for Zimmer Spine, a division of Zimmer, Inc. From September 2007 to October 2008, he served as the Senior Director, Marketing of Abbot Spine, Inc., from January 2007 to September 2007 as its Director, Product Management and from June 2005 to January 2007 he served as its Group Manager, Product Marketing. Mr. Whitfield received a B.S. degree in Business Administration, Marketing and Management from the University of Texas at Austin.

Board Composition

Our restated certificate of incorporation and restated bylaws provide that the authorized number of directors may be changed only by resolution of the board of directors. Seven directors are currently authorized. In accordance with our restated certificate of incorporation, our board of directors is divided into three classes with staggered three-year terms. At each annual meeting of stockholders, the successors to the directors whose terms then expire will be elected to serve until the third annual meeting following such election. Our directors are divided among the three classes as follows:

- The Class I directors are Max E. Link, Ph.D. and Jay M. Moyes and their terms will expire at the first annual meeting of stockholders to be held in 2015;
- The Class II directors are Eric K. Olson and David W. Truetzel, and their terms will expire at the second annual meeting of stockholders to be held in 2016; and
- The Class III directors are B. Sonny Bal, M.D. and Jeffrey S. White, and their terms will expire at the third annual meeting of stockholders to be held in 2017.

Any additional directorships resulting from an increase in the number of directors will be distributed among the three classes so that, as nearly as possible, each class will consist of one-third of the directors.

Section 16(a) Beneficial Ownership Reporting Compliance

As of December 31, 2013, our directors, officers and beneficial owners of more than ten percent of our common stock were not required to make period reports under Section 16 of the Exchange Act because our common stock had not been registered under Section 12 of the Exchange Act at such time.

Code of Business Conduct and Ethics

We have adopted a code of business conduct and ethics that applies to all of our employees, officers and directors, including those officers responsible for financial reporting. Our code of business conduct and ethics is available on our website at http://www.amedica.com/investors/corporate_governance. We intend to disclose any amendments to the code, or any waivers of its requirements on our website.

ITEM 11. EXECUTIVE COMPENSATION

The following discussion relates to the compensation of our “named executive officers,” including our Chief Executive Officer and President, Eric K. Olson, and our two most highly compensated executive officers (other than our chief executive officer), Jay M. Moyes, our Chief Financial Officer, and Bryan J. McEntire, our Chief Technology Officer.

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Summary Compensation Table

The following table sets forth information about certain compensation awarded or paid to our named executive officers for the 2012 and 2013 fiscal years.

<u>Name and Principal Position</u>	<u>Year</u>	<u>Salary (\$)</u>	<u>Bonus (\$)</u>	<u>Stock Awards \$(1)</u>	<u>Option Awards \$(2)</u>	<u>All Other Compensation \$(3)</u>	<u>Total (\$)</u>
Eric K. Olson	2013	315,192	—	442,000(4)	—	34,075(5)	791,267
Chief Executive Officer and President	2012	240,423	20,000	—	325,000	— (6)	585,423
Jay M. Moyes	2013	50,000(7)	100,000(8)	340,202(9)	—	30,288(10)	520,490
Chief Financial Officer	2012	—	—	—	—	8,043(11)	8,043
Bryan J. McEntire	2013	228,543	—	299,200(12)	—	33,246	560,989
Chief Technology Officer	2012	217,995	5,000	—	94,550	8,680	326,225

- (1) Amounts shown reflect the aggregate grant date fair value of restricted stock units, or RSUs, granted to the named executive officer computed in accordance with the Financial Accounting Standards Board, Accounting Standards Codification Topic 718, *Compensation — Stock Compensation*, or FASB ASC Topic 718. These amounts may not correspond to the actual value that will be recognized by the named executive officers. The grant date fair value of performance-based RSUs is determined based on the probable outcome of such performance conditions as of the grant date. The grant date fair value of the performance-based RSUs assuming the maximum potential value that can be achieved is \$442,000 for Mr. Olson, \$330,000 for Mr. Moyes and \$299,200 for Mr. McEntire. Assumptions used in the calculations of these amounts are included in Note 8 to our financial statements included in this Annual Report.
- (2) Amount shown for Mr. Olson reflects the grant date fair value of options awarded in 2012 determined in accordance with FASB ASC Topic 718. The amount shown for Mr. McEntire reflects the incremental fair value of stock options issued in exchange for outstanding stock options with exercise prices over \$25.77 per share in March 2012. These amounts exclude the value of estimated forfeitures. Assumptions used in the calculations of these amounts are included in Note 8 to our financial statements included elsewhere in this Annual Report.
- (3) Amount reflects the aggregation of any matching of 401(k) contributions, group term life premiums paid by us and accrued vacation in instances where each such amount is less than \$10,000, unless otherwise noted.
- (4) Amount includes the grant date fair value of (i) 23,279 RSUs issued to Mr. Olson in exchange for the cancellation of options to purchase 23,279 shares of our common stock held by Mr. Olson and (ii) 1,940 RSUs that were issued to Mr. Olson in June 2013.
- (5) Includes \$26,923 of accrued vacation.
- (6) Mr. Olson did not contribute money to our 401(k) plan in 2012. Therefore we paid no matching 401(k) amounts nor did we provide him with any other additional compensation in 2012.
- (7) Amount reflects the pro-rated amount of Mr. Moyes's annual salary of \$325,000 that was paid to Mr. Moyes since the beginning of his employment with us through the end of the 2013 fiscal year.
- (8) Amount reflects the signing bonus that was paid to Mr. Moyes upon the commencement of his employment.
- (9) Amount reflects the grant date fair value of (i) 58,197 RSUs that were issued to Mr. Moyes upon the commencement of his employment and (ii) 582 RSUs that were issued to Mr. Moyes as a non-employee director prior to the commencement of his employment.
- (10) Amount includes the \$27,185 of board member attendance fees paid to Mr. Moyes for his service as a member of our board of directors prior to the commencement of his employment with us.
- (11) Amount reflects fees paid to Mr. Moyes for service on our board of directors in the 2012 fiscal year.
- (12) Amount reflects the grant date fair value of (i) 16,101 RSUs that were issued to Mr. McEntire in February 2013 in exchange for the cancellation of options to purchase 16,101 shares of our common stock held by him and (ii) 970 RSUs that were issued to Mr. McEntire in June 2013.

Narrative Disclosure to Summary Compensation Table

Base Salaries. The base salaries for our named executive officers were determined by our compensation committee after reviewing a number of factors, including:

- the responsibilities associated with the position held by each of our executive officers and where that position fits within our overall corporate structure;
- the seniority of the individual executive's position;
- the base salary level of each executive officer in prior years;
- our overall financial position; and
- for executive officers other than our Chief Executive Officer, recommendations made by our Chief Executive Officer.

2012 Compensation

Our board of directors approved a salary of \$250,000 for Mr. Olson and \$228,545 for Mr. McEntire, effective as of January 1, 2012. Mr. Moyes was not employed by us at this time and as such received no compensation from us in 2012, other than fees for serving as a non-employee director. Our board of directors reduced the salaries of our named executive officers by ten percent effective as of May 31, 2012 in order to conserve cash. On December 12, 2012, Mr. McEntire's salary was restored to its original 2012 rate and Mr. Olson's salary was

increased to \$300,000. Our board, on the recommendation of our compensation committee

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decided to give this base salary increase to Mr. Olson in recognition of his significant contributions to our company, including helping to increase our revenue above \$23.0 million, aligning our sales and marketing team's salaries with our revenues and our receipt of 510(k) clearance for our second generation *Valeo* products while under his leadership. As a result of these salary changes in 2012, Messrs. Olson and McEntire received the following salary amounts:

<u>Name</u>	<u>Initial 2012 Base Salary (1)(\$)</u>	<u>10% Reduced 2012 Base Salary (2)</u>	<u>Restored/New 2012 Base Salary (3)</u>
Eric K. Olson Chief Executive Officer and President	250,000	225,000	300,000
Bryan J. McEntire Chief Technology Officer	228,545	205,691	228,545

- (1) Annual base salary rate from January 1, 2012 to May 31, 2012
(2) Annual base salary rate from June 1, 2012 to December 12, 2012
(3) Base salary as of December 12, 2012.

Annual Cash Bonuses. We have historically awarded discretionary cash bonuses to our executive officers. These bonuses are intended to reward our executive officers for the achievement of key strategic and business outcomes. Accordingly, each of Messrs. Olson and McEntire was awarded a cash bonus for 2012 equal to \$20,000 and \$5,000, respectively. Mr. Moyes was not employed with us in 2012 and as such did not receive a cash bonus for 2012. Our compensation committee has established a set of corporate objectives pursuant to which they may award our executive officers cash bonuses for their performance during 2013. These cash bonuses are discretionary and the compensation committee has not yet decided whether or not cash bonuses will be awarded for 2013 performance.

Long-Term Incentives. All options granted to our executive officers have been granted under the 2003 Stock Option Plan, or the 2003 Plan. These options vest over a period of time, generally four years. Upon termination of employment for any reason other than cause, our vested stock options granted to our named executive officers do not terminate and instead remain outstanding for their full-term of ten years. In the future, our compensation committee, with the approval of our board and stockholders, may grant to our named executive officers under the Amended and Restated 2012 Equity Incentive Plan, or the 2012 Plan, incentive stock options, non-qualified stock options, restricted and unrestricted stock awards, or stock-based awards, including RSUs and other stock based awards. See “—Equity Incentive Plans—2012 Plan” below for additional details about the 2012 Plan.

In March 2012, our board approved the cancellation of stock options held by current employees and members of our board with exercise prices above \$25.77 per share and replaced such options with new options for an equivalent number of shares with exercise prices of \$25.77 per share and ten-year terms to expiration, which were fully vested as of the date of grant. Mr. McEntire exchanged 8,147 options for an equal number of options at this time. None of Mr. Olson's options were cancelled because he held no options that had exercise prices over \$25.77. Mr. Moyes was not a member of our board of directors or our employee at that time.

2013 Compensation

In January 2013, we offered to each employee and director that held options to acquire shares of our common stock awarded under the 2003 Plan the opportunity to exchange such options for RSUs to be issued under the 2012 Plan on a one-for-one basis. As a result of the exchange offer, 93,968 RSUs were issued under the 2012 Plan in February 2013. Messrs. Olson and McEntire exchanged stock options and received 23,279 and 16,101 RSUs, respectively. Mr. Moyes did not have options eligible for conversion. The RSUs expire three years from the date of grant and will vest upon the earlier of (i) August 12, 2014 or (ii) the date of closing of a change in control provided, in each case, that the individual is providing services to us on such date.

In June 2013, in lieu of granting stock options, our board approved a grant of RSUs under the 2012 Plan to our named executive officers. Messrs. Olson and McEntire received 1,940 and 970 RSUs, respectively. The RSUs, which were awarded based on our 2012 performance, will vest upon the earlier of (i) August 12, 2014 or (ii) the date of a closing of a change in control, in each case, provided that the executive officer is providing services to us on such date and such event occurs within three years from the grant date.

On October 27, 2013, our board approved the hiring of Mr. Moyes as our Chief Financial Officer at an annual base salary of \$325,000 and provided Mr. Moyes with a signing bonus of \$100,000 which was paid on his first day of employment. In addition, if Mr. Moyes's employment is terminated by us for any reason other than cause, he shall be entitled to receive a pro-rated portion of his annual bonus for the year in which the termination occurs. In connection with his hiring, our board approved a grant of 58,197 RSUs to Mr. Moyes under the 2012 Plan, of which (i) 19,399 RSUs vested on his first day of employment, (ii) 19,399 RSUs vested on January 27, 2014 and (iii) 19,399 RSUs vested on February 12, 2014. In addition, the release date of the shares for any vested RSUs will be postponed until the first to occur of (i) a change in control or (ii) a separation of Mr. Moyes's service with us for any reason in compliance with Section 409A of the Internal Revenue Code.

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Our board authorized salary increases for each of our named executive officers effective as of October 30, 2013. Accordingly, Mr. Olson's salary was increased from \$300,000 to \$350,000, and Mr. McEntire's salary was increased from \$228,545 to \$235,000. Mr. Moyes did not receive a salary increase as he had just begun his employment with us at the time of salary increases.

2014 Compensation

On January 27, 2014, our board approved grants of RSUs to our named executive officers in the aggregate amount of 807,965 shares, as detailed below, subject to stockholder approval of an amendment to the 2012 Plan to increase the number of shares authorized for issuance under the 2012 Plan, which approval was obtained in February 2014, to be issued on the earlier of (i) a change in control or (ii) the effectiveness of a registration statement on Form S-8 which registers the shares underlying the RSUs:

<u>Name and Title</u>	<u>Grant Amount</u>
Eric K. Olson Chief Executive Officer and President	417,077
Jay M. Moyes Chief Financial Officer	299,713
Bryan J. McEntire Chief Technology Officer	91,175

Outstanding Equity Awards at Fiscal Year-End

The following table shows information regarding equity awards held by our named executive officers as of December 31, 2013:

<u>Name</u>	<u>Option Awards</u>				<u>Stock Awards</u>	
	<u>Number of Securities Underlying Unexercised Options (#) Exercisable</u>	<u>Number of Securities Underlying Unexercised Options (#) Unexercisable</u>	<u>Option Exercise Price (\$)</u>	<u>Option Expiration Date</u>	<u>Equity Incentive Plan Awards: Number of Unearned Shares, Units, or Other Rights That Have Not Vested (#)(1)</u>	<u>Equity Incentive Value of Unearned Market or Payout Value of Unearned Shares, Units or Other Rights That Have Not Vested (\$)(2)</u>
Eric K. Olson Chief Executive Officer and President	—	—	—	—	25,219	145,007
Jay M. Moyes Chief Financial Officer	—	—	—	—	39,380	226,434
Bryan J. McEntire Chief Technology Officer	7,760	—	6.44	6/8/2014	17,071	98,159

- (1) Mr. Olson and Mr. McEntire's RSUs vest upon the earlier of (i) August 12, 2014 or (ii) the date of a closing of a change in control provided, in each case, that the individual providing services to us on such date, and expire three years after the date of grant. 19,399 of Mr. Moyes RSUs vested on January 27, 2014 and 19,399 vested on February 12, 2014. The delivery of the vested RSUs will be postponed until the first to occur of (i) a change in control or (ii) a separation of Mr. Moyes service with us for any reason in compliance with Section 409A of the Internal Revenue Code.
- (2) Reflects the value calculated by multiplying the number of unvested RSUs by the value of our common stock on December 31, 2013, which we estimate was approximately \$5.75.

Retirement Benefits

401(k) Plan

We offer our executive officers, including our named executive officers, retirement benefits, including participation in our tax-qualified profit sharing plan that includes a "cash-or-deferred" (or 401(k)) feature in the same manner as other employees. The plan is intended to satisfy the requirements of Section 401 of the Internal Revenue Code. Our employees may elect to reduce their current compensation by up to the statutorily prescribed annual limit and have a like amount contributed to the plan. In addition, we may make discretionary and/or matching contributions to the plan in amounts determined annually by our board. We currently elect to

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match the contributions of our employees who participate in our 401(k) plan as follows: a match of 100% on the first 3% of compensation contributed by a plan participant and a match of 50% on amounts above 3%, up to 5%, of compensation contributed by a plan participant.

Potential Payments Upon Termination or Change in Control

We have entered into certain agreements and maintain certain plans that may require us to make certain payments and/or provide certain benefits to the executive officers named in the Summary Compensation Table in the event of a termination of employment or change in control.

Pursuant to severance agreements that we have entered into with each of our named executive officers (other than Mr. Moyes, who would receive severance payments pursuant to the terms of his employment agreement as more fully described below), upon the consummation of a change in control, all outstanding options, restricted stock and other such rights held by the executives will fully vest. Additionally, if a change in control occurs and at any time during the one-year period following the change in control (i) we or our successor terminate the executive's employment other than for cause (but not including termination due to the executive's death or disability) or (ii) the executive terminates his employment for good reason, then such executive has the right to receive payment consisting of a lump sum payment equal to two times his highest annual salary with us during the preceding three-year period, including the year of such termination and including bonus payments (measured on a fiscal year basis), but not including any reimbursements and amounts attributable to stock options and other non-cash compensation. "Change in control" is defined in the severance agreements as occurring upon: (i) any "person" (as such term is used in Sections 13(d) and 14(d) of the Exchange Act) becoming the "beneficial owner" (as defined in Rule 13d-3 under the Exchange Act), directly or indirectly, of securities representing 50% or more of the total voting power represented by our then outstanding voting securities (excluding securities held by us or our affiliates or any of our employee benefit plans) pursuant to a transaction or a series of related transactions which our board did not approve; (ii) a merger or consolidation of our company, other than a merger or consolidation which would result in our voting securities outstanding immediately prior thereto continuing to represent at least 50% of the total voting securities or such surviving entity or parent of such corporation outstanding immediately after such merger or consolidation; or (iii) the approval by our stockholders of an agreement for the sale or disposition of all or substantially all of our assets. As defined in the severance agreements, "cause" means: (i) the executive's commission of a felony (other than through vicarious liability or through a motor vehicle offense); (ii) the executive's material disloyalty or dishonesty to us; (iii) the commission by the executive of an act of fraud, embezzlement or misappropriation of funds; (iv) a material breach by the executive of any material provision of any agreement to which the executive and we are party, which breach is not cured within 30 days after our delivery to the executive of written notice of such breach; or (v) the executive's refusal to carry out a lawful written directive from our board. "Good reason" as defined in the severance agreements means, without the executive's consent: (i) a change in the principal location at which the executive performs his duties to a new work location that is at least 50 miles from the prior location; or (ii) a material change in the executive's compensation, authority, functions, duties or responsibilities, which would cause his position with us to become of less responsibility, importance or scope than his prior position, provided, however, that such material change is not in connection with the termination of the executive's employment with us for any reason.

In the event that an officer entitled to receive or receives payment or benefit under the severance agreements described above, or under any other plan, agreement or arrangement with us, or any person whose action results in a change in control or any other person affiliated with us and it is determined that the total amount of payments will be subject to excise tax under Section 4999 of the Internal Revenue Code, or any similar successor provisions, we will be obligated to pay such officer a "gross up" payment to cover all taxes, including any excise tax and any interest or penalties imposed with respect to such taxes due to such payment.

Pursuant to the terms of Mr. Moyes's employment arrangement, upon the occurrence of a change in control, all RSUs granted to Mr. Moyes at the time of his employment will fully vest. Pursuant to the employment arrangement, if: (i) Mr. Moyes is terminated by us without cause, (ii) he terminates his employment for good reason, or (iii) in the event a change in control occurs and Mr. Moyes is not offered continuing employment by the acquiring company or if such continuing employment is terminated without cause or if he terminates such continuing employment with good reason at any time during the twelve months following the change in control, we will be required to pay Mr. Moyes a lump sum equal to the sum of: (i) two times his annual salary in effect on the date of termination, (ii) any unpaid bonus through the end of his employment for the prior year that had been earned by Mr. Moyes but not paid plus a pro rata portion of his performance bonus and (iii) two times his sign on bonus. In addition, provided Mr. Moyes properly elects for continuation coverage, we will pay health insurance premiums for Mr. Moyes, his spouse and any covered dependents for a period of 24 months following the termination of his employment. Pursuant to the terms of the employment arrangement, "good reason" means, without Mr. Moyes's consent, the occurrence of any one or more of the following events: (i) a material diminution of Mr. Moyes's authority, functions, duties or responsibilities; (ii) a relocation of Mr. Moyes's principal workplace to a new location more than 50 miles from the prior location; (iii) the material diminution of Mr. Moyes's annual base salary, other than in the event of a reduction in compensation of all of our executive officers, generally, so long as the reduction to Mr. Moyes's base salary is no more than the average reduction; or (iv) a material breach by us of the terms of the employment agreement. "Change in control" has the same meaning as previously described under the severance agreements for the other executive officers.

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In connection with a change in control, if Mr. Moyes will be required to pay any excise tax pursuant to Section 4999 of the Internal Revenue Code or any similar successor provisions then we will make an additional “gross up” payment to Mr. Moyes in an amount to cover such excise tax and the taxes associated with each payment.

Director Compensation

The following table shows the total compensation paid or accrued during the fiscal year ended December 31, 2013 to each of our non-employee directors except Mr. Jeffrey S. White who was not a member of our board of directors in 2013. Fees paid to Mr. Moyes for his service as a director prior to his employment with us are included in the Summary Compensation Table.

<u>Name</u>	<u>Fees Earned or Paid in Cash(\$)</u>	<u>Stock Awards (\$)(1)(2)</u>	<u>Option Grants (\$)(3)</u>	<u>All Other Compensation (\$)</u>	<u>Total (\$)</u>
Max E. Link, Ph.D.	38,500	10,200	—	—	48,700
B. Sonny Bal, M.D.	26,000	10,200	—	—	36,200
Gregg R. Honigblum(4)	24,151	10,200	—	—	34,351
Rohit Patel(5)	30,909	10,200	—	12,250(6)	53,359
George Singer(7)	17,522	10,200	—	—	27,722
David Truetzel	43,751	10,200	—	—	53,951

- (1) As part of the annual grant to directors, each director received 582 RSUs. These RSUs expire three years from the date of grant and will only vest upon the earlier of (a) the date of expiration of the lock-up period imposed in connection with the closing of an underwritten initial public offering of shares of our common stock or (b) the date of a closing of a change of control provided, in either case, that the applicable vesting event occurs within three years following the date of grant.
- (2) Amount shown reflects the grant date fair value of the RSUs awarded in 2013 determined in accordance with the Financial Accounting Standards Board, Accounting Standards Codification Topic 718, *Compensation-Stock Compensation*. Assumptions used in the calculation of these amounts are included in Note 8 to our financial statements included elsewhere in this Annual Report.
- (3) No stock options were granted to directors during 2013. However, as of December 31, 2013, our directors held the following aggregate number of stock options: Dr. Link, 3,783; Dr. Bal, 2,813; and Mr. Honigblum, 4,171. Mr. Singer, did not hold any stock options as of December 31, 2013. All stock options are fully vested.
- (4) Mr. Honigblum resigned from our board of directors in September 2013.
- (5) Mr. Patel resigned from our board of directors in September 2013.
- (6) We paid Mr. Patel \$12,250 in consulting fees related to a litigation matter against our former Chief Executive Officer, Ben Shappley, in 2013.
- (7) Mr. Singer resigned from our board of directors in September 2013.

We compensate each of the non-employee members of our Board in accordance with the following annual retainer and meeting fees (paid on a quarterly basis):

• Board member Annual Retainer	\$20,000
• Board Chair Annual Additional Retainer	\$10,000
• Committee Chair Annual Retainer	\$ 7,500
• Committee member Annual Retainer	\$ 3,750
• Board meeting-in person attendance	\$ 1,500
• Board meeting-telephonic attendance	\$ 1,000
• Committee meeting attendance	\$ 1,500
• Committee meeting-telephonic attendance	\$ 1,000

In addition to cash compensation, the non-employee members of our Board have historically been awarded an annual stock option grant in the amount of 582 shares of our common stock. In 2013, in lieu of stock options, non-employee members of the board were each granted 582 RSUs. In January 2013, we offered to each director that held options to acquire shares awarded under the 2003 Plan the opportunity to exchange such options for an equal number of RSUs. Messrs. Patel and Truetzel each exchanged stock options and received 9,021 RSUs and 2,716 RSUs, respectively, under the 2012 Plan. These RSUs expire three years from the date of grant and will only vest upon continued service with us and if either of the following events occurs prior to the expiration date: (i) August 12, 2014, the date of the expiration of the lock-up period imposed on the directors after the completion of the closing of the initial public offering of the shares of our common stock or (ii) upon a “change in control”. “Change of Control” is defined in the restricted stock unit agreements entered into in connection with the awarding of the RSUs, as occurring upon: (i) any “person” (as such term is used in Sections 13(d) and 14(d) of the Exchange Act) becoming the “beneficial owner” (as defined in Rule 13d-3 under the Exchange Act), directly or indirectly, of securities representing 50% or more of the total voting power represented by our then outstanding voting securities (excluding for this purpose any such voting securities held by the us our affiliates or by any of our employee benefit plans) pursuant to a transaction or a series of related transactions which the Board of Directors does not approve; or (ii) a merger or consolidation of our company, whether or not approved by the Board of Directors, other than a merger or consolidation which would result in our voting securities outstanding immediately prior thereto continuing to represent (either by remaining outstanding or by being converted into voting securities of the surviving entity or the parent of such corporation) more than 50% of the total voting power represented by our voting securities of the surviving entity or parent of such corporation, as the case may be, outstanding immediately after such merger or consolidation; or (iii) the sale or disposition by us of all or substantially all of our assets in a transaction requiring stockholder approval.

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

The following table sets forth certain information regarding the beneficial ownership of our common stock as of March 14, 2014 by:

- each of our current directors;
- the executive officers named in the summary compensation table;
- all of our directors and executive officers as a group; and
- each stockholder known by us to own beneficially more than 5% of our common stock.

Beneficial ownership is determined in accordance with the rules of the SEC and includes voting or investment power with respect to the securities. Shares of common stock that may be acquired by an individual or group within 60 days of March 14, 2014, pursuant to the exercise of options or warrants, are deemed to be outstanding for the purpose of computing the percentage ownership of such individual or group, but are not deemed to be outstanding for the purpose of computing the percentage ownership of any other person shown in the table. Percentage of shares beneficially owned is based on 12,310,354 shares issued and outstanding on March 14, 2014.

Except as indicated in footnotes to this table, we believe that the stockholders named in this table have sole voting and investment power with respect to all shares of common stock shown to be beneficially owned by them, based on information provided to us by such stockholders. The address for each director and executive officer listed is: c/o Amedica Corporation, 1885 West 2100 South, Salt Lake City, Utah 84119.

<u>Name and Address of Beneficial Owner</u>	<u>Shares Beneficially Owned</u>	
	<u>Number</u>	<u>Percentage</u>
Directors and Named Executive Officers:		
Max E. Link, Ph.D.(1)	76,291	*%
B. Sonny Bal, M.D.(2)	57,298	*
David W. Truetzel(3)	33,239	*
Jeffrey S. White	—	*
Jay M. Moyes(4)	66,465	*
Eric K. Olson	—	*
Bryan J. McEntire(5)	7,759	*
All directors and executive officers as a group (12 individuals)(6)	241,052	1.9%
Five Percent Stockholders:		
Karl Kipke(7)	1,575,012	11.4%
Hampshire Group, LLC 500 Plaza on the Lake, Suite #103 Austin, TX 78746		

* Represents beneficial ownership of less than 1% of the shares of our common stock.

- (1) Consists of 70,085 shares of our common stock, options to acquire 3,782 shares of our common stock currently exercisable or exercisable within 60 days of March 14, 2014. Also includes 2,424 common stock warrants that are currently exercisable.
- (2) Consists of 18,750 shares of our common stock held by Dr. Bal, 33,894 shares of our common stock held by Dr. Bal and his spouse, options to acquire 3,200 shares of our common stock currently exercisable or exercisable within 60 days of March 14, 2014. Also includes 1,454 common stock warrants that are currently exercisable.
- (3) Consists of 337 shares of our common stock held by Mr. Truetzel, 50% of 22,129 shares of our common stock held by Spinal Management, LLC, of which Mr. Truetzel is a 50% member, 15,625 shares of our common stock held by Truetzel Revocable Trust of which Mr. Truetzel and his spouse are the sole beneficiaries. Also includes 1,212 common stock warrants that are currently exercisable.
- (4) Consists of 58,197 RSUs, 1,534 shares of our common stock, 6,250 shares of our common stock that are beneficially owned by Drayton Investments, LLC, and 484 common stock warrants that are immediately exercisable and beneficially owned by Drayton Investments, LLC. Mr. Moyes is a managing member of Drayton Investments, LLC.
- (5) Consists of options to acquire 7,759 shares of our common stock currently exercisable or exercisable within 60 days of March 14, 2014, and does not include 17,071 RSUs.

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- (6) Consists of 176,592 shares of our common stock, options to acquire 14,741 shares of our common stock, and 5,574 common stock warrants that are currently exercisable.
- (7) Consists of: (i) 16,179 warrants that are currently exercisable held by Mr. Kipke; (ii) 1,303,356 shares and 96,994 common stock warrants that are currently exercisable held by Hampshire Med Tech Partners, LP; (iii) 25,908 shares held by Hampshire Healthcare Partners, LP; and (iv) 16,919 shares held by Hampshire Asset Management, LLC. Hampshire Med Tech is the general partner of Hampshire Med Tech Partners, LP and Special Opportunities is the general partner of Hampshire Healthcare Partners, LP. Mr. Kipke is the managing member of each of Hampshire Med Tech and Special Opportunities and the president of Hampshire Asset Management, LLC. Also includes 115,655 shares held by KM Healthcare Holdings, LP. No Footprints is the general partner of KM Healthcare Holdings, LP. Mr. Kipke is a managing member of No Footprints and shares voting and dispositive power with Mr. Murphy with respect to the shares held by KM Healthcare Holdings, LP.

Equity Compensation Plan Information

The following table summarizes our equity compensation plans as of December 31, 2013:

<u>Plan category</u>	<u>(a) Number of securities to be issued upon exercise of outstanding options, warrants, and rights</u>	<u>(b) Weighted-average exercise price of outstanding options, warrants and rights</u>	<u>(c) Number of securities remaining available for future issuances under equity compensation plans (excluding securities reflected in column (a))</u>
Equity compensation plans approved by security holders	294,775	\$ 28.90	91,659
Equity compensation plans not approved by security holders	—	\$ 0.00	N/A
Total	294,775	\$ 28.90	91,659

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

The following includes a summary of transactions since January 1, 2012 to which we have been a party, in which the amount involved in the transaction exceeded \$120,000, and in which any of our directors, executive officers or, to our knowledge, beneficial owners of more than 5% of our common stock, on an as converted basis, or any member of the immediate family of any of the foregoing persons had or will have a direct or indirect material interest, other than equity and other compensation, termination, change in control and other arrangements, which are described under "Executive and Director Compensation." With the approval of our board of directors, we have engaged in the transactions described below with our directors, executive officers and beneficial owners of more than 5% of our common stock, on an as converted basis, and affiliates of our directors, executive officers and 5% stockholders.

Transactions with MSK Investments, LLC and its Affiliates following our Acquisition of US Spine, Inc.

Settlement Agreement with MSK Investments, LLC

In September 2010, we acquired US Spine, Inc., or US Spine. In this transaction, US Spine became our wholly owned subsidiary as we acquired all of the outstanding capital stock of US Spine. In May 2012, we entered into a settlement agreement with James G. Koman and with MSK Investments, LLC, or MSK, a company controlled by Mr. Koman, on its own behalf and acting in its capacity as stockholders' representative for the former stockholders of US Spine, to resolve certain disputes. Pursuant to the settlement agreement, in lieu of the issuance of up to 6,250,000 shares of our Series E convertible preferred stock upon the achievement of certain earnout milestones, which we refer to as the US Spine Earnout, we issued (a) 842,443 shares of our Series E convertible preferred stock to the former stockholders of US Spine, of which 39,249 shares were issued to MSK and its affiliates, and (b) 2,557,562 shares of our Series C convertible preferred stock to MSK. We also agreed to release the 1,806,250 shares of our Series E convertible preferred stock from an escrow arrangement established at the time of our acquisition of US Spine, of which 1,380,654 were received by MSK and its affiliates. MSK and Mr. Koman also agreed to certain standstill covenants in our favor that expire on May 10, 2015. Spinal Management LLC, of which David Truetzel is a 50% co-owner, also received a commission that was paid in 42,122 and 127,878 of the shares of our Series E convertible preferred stock and Series C convertible preferred stock, respectively, issued under the settlement agreement.

Restructuring and Payment of the US Spine Note

In October 2012, we restructured the terms of a promissory note issued in favor of MSK at the time of our acquisition of US Spine, or the US Spine Note, to extend the maturity date of the second \$3.0 million installment from September 2012 to December 2012. We made payments to MSK of \$500,000 on October 31, 2012 and \$2,500,000 on December 17, 2012 in connection with this restructuring.

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Restructuring and Conversion of Senior Secured Subordinated 6%/8% Convertible Promissory Notes

Between March 2011 and May 2011, we issued an aggregate principal amount of \$24.8 million of Senior Secured Subordinated 6%/8% Convertible Promissory Notes, or the Senior Secured Notes, and warrants to purchase 288,685 shares of our common stock at an exercise price of \$51.55 per share to 85 accredited investors. In connection with the initial closing of this offering, we received a commitment from Hampshire Med Tech Partners, LP to purchase an additional \$5.0 million Senior Secured Note by no later than the first anniversary of the initial closing of that offering. Pursuant to this commitment, we issued an additional \$5.0 million Senior Secured Note to Hampshire Med Tech Partners, LP in February 2012.

In December 2012, we amended the terms of the Senior Secured Notes and the holders thereof converted all of their Senior Secured Notes into an aggregate of 14,887,500 shares of our Series F convertible preferred stock. We also amended the terms of the warrants to purchase shares of our common stock issued in connection with the issuance of the Senior Secured Notes to lower the exercise prices thereof from \$51.55 per share to \$25.77 per share. As a result, we issued an aggregate of 6,131,250 shares of our Series F convertible preferred stock to the following directors, officers and beneficial owners of more than 5% of our common stock, on an as converted basis, and their affiliates:

<u>Name</u>	<u>Number of Shares of Series F Convertible Preferred Stock</u>
Max E. Link, Ph.D.	25,000
David Truetzel(1)	12,500
Allan R. Lyons(2)	475,000
Gregg R. Honigblum(3)	6,250
Karl Kipke(4)	5,000,000
B. Sonny Bal, M.D.(5)	12,500
Kevin Murphy	600,000

- (1) Includes 12,500 shares that were issued to Truetzel Revocable Trust, of which Mr. Truetzel and his spouse are the sole beneficiaries.
- (2) Shares were issued to Vestal. Mr. Lyons is the managing member and sole owner of 21st Century Strategic Investment Planning, LLC, the general partner of Vestal.
- (3) Represents 50% of the 12,500 shares that were issued to Creation Capital. Mr. Honigblum is a 50% owner and a managing member of Creation Capital. Mr. Honigblum resigned from our board of directors in September 2013.
- (4) Shares were issued to Hampshire Med Tech Partners, LP. Mr. Kipke is the managing member of Hampshire Med Tech, its general partner.
- (5) Shares were issued to Dr. Bal's father.

Warrant Restructuring and Private Placement of Common Stock

In March 2013, we amended the terms of certain of the common stock warrants issued in connection with the issuance of the Senior Secured Notes to further lower the exercise prices thereof from \$25.77 per share to \$17.53 per share. We then issued an aggregate of 178,516 shares of our common stock to 33 accredited investors upon exercise of the amended common stock warrants and the sale of additional shares of our common stock to other investors in the offering at \$17.53 per share. We also issued to investors who exercised their common stock warrants new warrants to purchase an aggregate of 76,455 shares of our common stock at an exercise price of \$17.53 per share. We issued an aggregate of 53,347 shares of our common stock and new warrants to purchase up to 17,773 shares of our common stock at an exercise price of \$17.53 per share to the following directors, officers and beneficial owners of more than 5% of our common stock, on an as converted basis, and their affiliates:

<u>Name</u>	<u>Common Stock upon Exercise of Warrants</u>	<u>New Common Stock</u>	<u>New Common Stock Warrants</u>
Allan R. Lyons(1)	9,214	—	9,214
Kevin Murphy	8,558	—	8,558
Karl Kipke(2)	—	53,347	—

- (1) Represents the exercise of common stock warrants by, and issuance of common stock warrants to, Vestal. Mr. Lyons is the managing member and sole owner of 21st Century Strategic Investment Planning, LLC, the general partner of Vestal.
- (2) Represents 53,347 shares of common stock purchased by Hampshire Med Tech Partners, LP. Mr. Kipke is the managing member of Hampshire Med Tech, its general partner.

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Private Placement of Series F Convertible Preferred Stock

In August 2013 and September 2013, we issued an aggregate of 94.8 units, each unit consisting of 50,000 shares of our Series F convertible preferred stock and a warrant to acquire 970 shares of our common stock at an exercise price of \$25.77 per share, to 45 accredited investors at \$100,000 per unit. This resulted in our issuance of an aggregate of 4,740,000 shares of our Series F convertible preferred stock and warrants to purchase an aggregate of 91,951 shares of our common stock, including an aggregate of 1,125,000 shares of our Series F convertible preferred stock and warrants to purchase an aggregate of 21,824 shares of our common stock to the following directors, officers and beneficial owners of more than 5% of our common stock, on an as converted basis, and their affiliates:

<u>Name</u>	<u>Number of Units</u>	<u>Purchase Price</u>	<u>Number of Shares of Series F Convertible Preferred Stock</u>	<u>Common Stock Warrants</u>
Max E. Link, Ph.D.	2.0	\$ 200,000	100,000	1,940
B. Sonny Bal, M.D.	1.5	\$ 150,000	75,000	1,455
David W. Truetzel(1)	1.0	\$ 100,000	50,000	970
Jay M. Moyes(2)	0.5	\$ 50,000	25,000	485
George Singer(3)	1.0	\$ 100,000	50,000	970
Allan R. Lyons(4)	3.5	\$ 350,000	175,000	3,395
James G. Koman(5)	1.0	\$ 100,000	50,000	970
Kevin Murphy(6)	12.0	\$ 1,200,000	600,000	11,639

- (1) Investment made by Truetzel Revocable Trust, of which Mr. Truetzel and his spouse are the sole beneficiaries.
- (2) Investment made by Drayton Investments, LLC, of which Mr. Moyes is a managing member.
- (3) Consists of 50% of the investment made by Singer Bros. LLC. Mr. Singer is a 50% owner and a managing member of Singer Bros. LLC. Mr. Singer resigned from our board of directors in September 2013.
- (4) Investment made by Vestal. Mr. Lyons is the managing member and sole owner of 21st Century Strategic Investment Planning, LLC, the general partner Vestal.
- (5) Investment made by MSK, of which Mr. Koman is the managing member.
- (6) In connection with the sale and issuance of certain of the units in this financing, we also issued to TGP Securities, Inc., an entity controlled by Mr. Murphy, warrants to purchase 9,311 shares of our common stock at an exercise price of \$56.70 per share and paid a cash commission of \$480,000 to TGP Securities, Inc., neither of which are reflected in the table.

Transactions with Creation Capital, LLC and Creation Capital Advisors, LLC

Mr. Gregg R. Honigblum, the Chief Executive Officer and a 50% co-owner of each of Creation Capital LLC, or Creation Capital, and Creation Capital Advisors, LLC, or Creation Advisors, served on our board of directors from 2006 until September 2013. In connection with the private placement of our Senior Secured Notes Creation Capital served as our placement agent In February 2012, when we issued an additional \$5.0 million Senior Secured Note to Hampshire Med Tech Partners, LP, we paid Creation Capital an \$212,500 as commissions.

In June 2012, we entered into a financial advisor consulting agreement with Creation Advisors, pursuant to which we agreed to extend the termination date of certain Series C convertible preferred stock warrants previously issued to Creation Capital from February 2013 to February 2018.

In connection with the conversion of our Senior Secured Notes in December 2012, we agreed to pay Creation Advisors a strategic financial advisory fee in the amount of approximately \$447,000. We agreed to pay half of the advisory fee, approximately \$223,000 in December 2012 and the remaining half within 24 months, which we paid in September 2013. Karl Kipke, who beneficially owns more than 5% of our common stock, received \$60,000 from Creation Advisors in 2012, as a consultant for Creation Advisors, for advising us at this time on our financing options.

In connection with the warrant restructuring and private placement of common stock in March 2013, we paid Creation Advisors a strategic financial advisory fee of approximately \$250,000. In October 2013, we entered into a one-year consulting agreement for financial advisory services with Creation Advisors in which Creation Advisors will receive compensation of up to \$180,000 in cash (payable \$15,000 per month). We paid \$45,000, under this agreement, during the year ended December 31, 2013.

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Director Independence

Our board of directors has reviewed the materiality of any relationship between us and each of our directors, either directly or indirectly. Based on this review, the board of directors has determined that Max E. Link, Ph.D., David W. Truetzel, Jeffrey S. White and B. Sonny Bal, M.D. are “independent directors” as defined by the SEC and NASDAQ. The rules of The NASDAQ Capital Market require that a majority of the board of directors of a listed company consist of independent directors, as defined by the rules of The NASDAQ Capital Market. We currently have a board of directors consisting of a majority of independent directors.

Policy for Approval of Related Person Transactions

We believe that all the transactions described above were made on terms no less favorable to us than those that could have been obtained from unaffiliated third parties. With the exception of transactions in which related parties participated on the same terms as those of other participants who were not related parties, our board of directors reviewed and approved the transactions with each related party, namely our directors, executive officers and beneficial owners of more than 5% of our common stock, on an as converted basis, and affiliates of our directors, executive officers and 5% stockholders, and reviewed the material facts as to a related party’s relationship or interest in a transaction that were disclosed to our board of directors prior to our board of directors’ consideration of a transaction with a related party. The transactions involving related parties were approved by our board of directors, including all of our directors who were not interested in these transactions.

All future related party transactions will be approved by our audit committee. Pursuant to the written charter of our audit committee, the audit committee is responsible for reviewing and approving, prior to our entry into any transaction involving related parties, all transactions in which we are a participant and in which any parties related to us has or will have a direct or indirect material interest.

In reviewing and approving these transactions, the audit committee shall obtain, or shall direct our management to obtain on its behalf, all information that the committee believes to be relevant and important to a review of the transaction prior to its approval. Following receipt of the necessary information, a discussion shall be held of the relevant factors, if deemed to be necessary by the committee, prior to approval. If a discussion is not deemed to be necessary, approval may be given by written consent of the committee. No related party transaction shall be entered into prior to the completion of these procedures.

The audit committee or its chairman, as the case may be, shall approve only those related party transactions that are determined to be in, or not inconsistent with, the best interests of us and our stockholders, taking into account all available facts and circumstances as the committee or the chairman determines in good faith to be necessary. No member of the audit committee shall participate in any review, consideration or approval of any related party transaction with respect to which the member or any of his or her immediate family members is the related party.

ITEM 14. PRINCIPAL ACCOUNTANT FEES AND SERVICES

The following table represents aggregate fees billed to us for the years ended December 31, 2013 and 2012 by Ernst & Young LLP. All fees below were approved by our Audit Committee.

	Year Ended December 31,	
	2013	2012
Audit fees (1)	\$1,187,300	\$265,372
Audit related fees	—	—
Tax fees	—	—
All other fees	—	—
	<u>\$1,187,300</u>	<u>\$265,372</u>

- (1) Audit fees consist of fees incurred for professional services rendered for the audit of our annual financial statements and review of the quarterly financial statements that are normally provided by Ernst & Young LLP in connection with regulatory filings or engagements. For the year ended December 31, 2013, audit fees also includes fees related to our initial public offering and review of documents filed with the SEC.

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Pre-approval Policies and Procedures

Our Audit Committee has responsibility for establishing policies and procedures for the pre-approval of audit and non-audit services rendered by our independent registered public accounting firm, Ernst & Young LLP. The policy generally pre-approves specified services in the defined categories of audit services, audit-related services, and tax services up to specified amounts. Pre-approval may also be given as part of the Audit Committee's approval of the scope of the engagement of the independent registered public accounting firm or on an individual explicit case-by-case basis before the independent registered public accounting firm is engaged to provide each service. The pre-approval of services may be delegated to one or more of the Audit Committee's members, but the decision must be reported to the full Audit Committee at its next scheduled meeting.

The Audit Committee has determined that the rendering of the above services by Ernst & Young LLP is compatible with the SEC's policies on auditor independence.

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PART IV

ITEM 15. EXHIBITS AND FINANCIAL STATEMENT SCHEDULES

Reference is made to the Index to Consolidated Financial Statements beginning on Page F-1 hereof.

(1) *Financial Statements*. The following consolidated financial statements and the notes thereto, and the Report of Independent Registered Public Accounting Firms are incorporated by reference as provided in Item 8 of this report:

• Report of Independent Registered Public Accounting Firm	F-2
• Consolidated Balance Sheets at December 31, 2013 and 2012	F-3
• Consolidated Statements of Operations and Comprehensive Loss for the Years Ended December 31, 2013 and 2012	F-4
• Consolidated Statements of Convertible Preferred Stock and Stockholders' Deficit for the Period from December 31, 2011 through December 31, 2013	F-5
• Consolidated Statements of Cash Flows for the Years Ended December 31, 2013 and 2012	F-6
• Notes to Consolidated Financial Statements	F-7

(2) Consolidated Financial Statement Schedules

Consolidated Financial Statement Schedules have been omitted because they are either not required or not applicable, or because the information required to be presented is included in the consolidated financial statements or the notes thereto included in this Annual Report.

(3) Exhibits

The exhibits listed on the accompanying Exhibit Index are filed or incorporated by reference as part of this Annual Report and such Exhibit Index is incorporated by reference.

<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
3.1	Restated Certificate of Incorporation of the Registrant		Form 8-K (Exhibit 3.1)	2/20/14	001-33624
3.2	Restated Bylaws of the Registrant		Form 8-K (Exhibit 3.1)	2/20/14	001-33624
4.1	Form of Common Stock Certificate of the Registrant		Amendment No. 3 to Form S-1 (Exhibit 4.1)	1/29/14	333-192232
4.2	Fifth Amended and Restated Registration Rights Agreement by and among the Registrant and certain of its stockholders, dated as of July 27, 2010		Form S-1 (Exhibit 4.2)	11/8/13	333-192232
4.3	Warrant Agreement by and between the Registrant and Creation Capital LLC, dated as of February 24, 2006		Form S-1 (Exhibit 4.5)	11/8/13	333-192232

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
4.4	Series D Warrant Agreement by and between the Registrant and Creation Capital LLC, dated as of April 27, 2007		Form S-1 (Exhibit 4.6)	11/8/13	333-192232
4.5	Common Stock Warrant Agreement by and between the Registrant and Creation Capital LLC, dated as of April 30, 2008		Form S-1 (Exhibit 4.7)	11/8/13	333-192232
4.6	Series E Warrant Agreement by and between the Registrant and Creation Capital LLC, dated as of September 14, 2010		Form S-1 (Exhibit 4.8)	11/8/13	333-192232
4.7	Form of Warrant to Purchase Shares of Common Stock of the Registrant, issued on May 9, 2011		Amendment No. 3 to Form S-1 (Exhibit 4.9)	1/29/14	333-192232
4.8	Warrant to Purchase 156,978 Shares of Series F Convertible Preferred Stock by and between the Registrant and GE Capital Equity Investments, Inc., dated as of December 17, 2012		Form S-1 (Exhibit 4.10)	11/8/13	333-192232
4.9	Warrant to Purchase 113,022 Shares of Series F Convertible Preferred Stock by and between the Registrant and Zions First National Bank, dated as of December 17, 2012		Form S-1 (Exhibit 4.11)	11/8/13	333-192232
4.10	Form of Warrant to Purchase Shares of Common Stock of the Registrant, issued on March 4, 2011 and May 9, 2011		Form S-1 (Exhibit 4.12)	11/8/13	333-192232
4.11	Form of Amendment to Warrant to Purchase Shares of Common Stock of the Registrant, dated as of December 18, 2012		Form S-1 (Exhibit 4.13)	11/8/13	333-192232

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
4.12	Form of Amendment No. 2 to Warrant to Purchase Shares of Common Stock of the Registrant, dated as of February 1, 2013		Form S-1 (Exhibit 4.14)	11/8/13	333-192232
4.13	Warrant to Purchase Shares of Common Stock of the Registrant by and between the Registrant and the University of Utah Research Foundation, dated as of February 17, 2010		Form S-1 (Exhibit 4.15)	11/8/13	333-192232
4.14	Form of Warrant to Purchase Shares of Common Stock of the Registrant, issued on April 18, 2011, November 15, 2011, November 16, 2011, February 22, 2012, February 29, 2012 and March 7, 2012		Form S-1 (Exhibit 4.16)	11/8/13	333-192232
4.15	Form of Warrant to Purchase Shares of Common Stock of the Registrant, issued on August 30, 2013 and September 20, 2013, as amended		Amendment No. 2 to Form S-1 (Exhibit 4.17)	12/20/13	333-192232
4.16	Form of Amendment to Warrant to Purchase Common Stock of the Registrant, dated as of December 23, 2013		Amendment No. 3 to Form S-1 (Exhibit 4.17.1)	1/29/14	333-192232
4.17	Warrant to Purchase Shares of Common Stock of the Registrant by and between the Registrant and Zions First National Bank, dated as of March 17, 2011		Form S-1 (Exhibit 4.18)	11/8/13	333-192232
4.18	Series E Warrant Agreement by and between the Registrant and Zions First National Bank, dated as of April 7, 2010		Amendment No. 2 to Form S-1 (Exhibit 4.19)	12/20/13	333-192232

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
4.19	Form of Warrant to Purchase Shares of Common Stock of the Registrant, issued to TGP Securities, Inc. on August 30, 2013 and September 20, 2013, as amended		Amendment No. 2 to Form S-1 (Exhibit 4.20)	12/20/13	333-192232
4.20	Form of Amendment to Warrant to Purchase Shares of Common Stock of the Registrant, issued to TGP Securities, Inc., dated as of December 23, 2013		Amendment No. 3 to Form S-1 (Exhibit 4.21)	1/29/14	333-192232
10.1	Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and the financial institutions party thereto, dated as of December 17, 2012		Form S-1 (Exhibit 10.1)	11/8/13	333-192232
10.2	First Amendment to Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and US Spine, Inc., dated as of June 28, 2013		Form S-1 (Exhibit 10.2)	11/8/13	333-192232
10.3	Second Amendment and Waiver to Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and US Spine, Inc., dated as of July 31, 2013		Form S-1 (Exhibit 10.3)	11/8/13	333-192232
10.4	Third Amendment and Waiver to Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and US Spine, Inc., dated as of August 15, 2013		Form S-1 (Exhibit 10.4)	11/8/13	333-192232
10.5	Fourth Amendment to Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and US Spine, Inc., dated as of December 23, 2013		Amendment No. 3 to Form S-1 (Exhibit 10.4.1)	1/29/14	333-192232

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
10.6	Fifth Amendment to Loan and Security Agreement by and among the Registrant, General Electric Capital Corporation and US Spine, Inc., dated as of January 28, 2014.		Amendment No. 3 to Form S-1 (Exhibit 10.4.2)	1/29/14	333-192232
10.7	Pledge Agreement by and between the Registrant and General Electric Capital Corporation, dated as of December 17, 2012		Form S-1 (Exhibit 10.5)	11/8/13	333-192232
10.8	Intellectual Property Security Agreement by the Registrant in favor of General Electric Capital Corporation, dated as of December 17, 2012		Form S-1 (Exhibit 10.6)	11/8/13	333-192232
10.9	Joint Development and License Agreement by and between the Registrant and Orthopaedic Synergy, Inc., dated as of February 8, 2010**		Amendment No. 3 to Form S-1 (Exhibit 10.7)	1/29/14	333-192232
10.10	Distribution Agreement by and between the Registrant and Orthopaedic Synergy, Inc., dated as of February 22, 2010, and First Amendment and Addendum thereto, dated as of November 1, 2012**		Amendment No. 3 to Form S-1 (Exhibit 10.8)	1/29/14	333-192232
10.11	Centrepointe Business Park Lease Agreement Net by and between the Registrant and Centrepointe Properties, LLC, dated as of April 21, 2009		Form S-1 (Exhibit 10.10)	11/8/13	333-192232
10.12	First Addendum to Centrepointe Business Park Lease Agreement Net by and between the Registrant and Centrepointe Properties, LLC, dated as of January 31, 2012		Form S-1 (Exhibit 10.11)	11/8/13	333-192232
10.13	Form of Severance and Change of Control Agreement*		Amendment No. 2 to Form S-1 (Exhibit 10.12)	12/20/13	333-192232

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
10.14	Employment Term Sheet by and between the Registrant and Jay M. Moyes, dated as of October 29, 2013*		Amendment No. 2 to Form S-1 (Exhibit 10.13)	12/20/13	333-192232
10.15	Restricted Stock Unit Agreement, by and between the Registrant and Jay Moyes, dated as of October 30, 2013*		Amendment No. 3 to Form S-1 (Exhibit 10.13.1)	1/29/14	333-192232
10.16	Form of Indemnification Agreement by and between the Registrant and its officers and directors 8		Amendment No. 2 to Form S-1 (Exhibit 10.14)	12/20/13	333-192232
10.17	Amedica Corporation Amended and Restated 2012 Equity Incentive Plan*		Amendment No. 4 to Form S-1 (Exhibit 10.15)	2/12/14	333-192232
10.18	Form of 2012 Stock Option Grant Notice and Stock Option Agreement*		Amendment No. 4 to Form S-1 (Exhibit 10.16)	2/12/14	333-192232
10.19	Form of 2012 Restricted Stock Award and Restricted Stock Unit Agreement*		Amendment No. 4 to Form S-1 (Exhibit 10.17)	2/12/14	333-192232
10.20	Amedica Corporation 2003 Stock Option Plan*		Form S-1 (Exhibit 10.18)	11/8/13	333-192232
10.21	Form of 2003 Non-Qualified Stock Option Agreement and Notice of Exercise of Non-Qualified Stock Option thereunder*		Form S-1 (Exhibit 10.19)	11/8/13	333-192232
10.22	Form of 2003 Incentive Stock Option Agreement and Notice of Exercise of Incentive Stock Option thereunder*		Form S-1 (Exhibit 10.20)	11/8/13	333-192232
21.1	List of Subsidiaries of the Registrant		Form S-1 (Exhibit 21.1)	11/8/13	333-192232
31.1	Certification of Chief Executive Officer	X			
31.2	Certification of Chief Financial Officer	X			

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<u>Exhibit Number</u>	<u>Exhibit Description</u>	<u>Filed with this Report</u>	<u>Incorporated by Reference herein from Form or Schedule</u>	<u>Filing Date</u>	<u>SEC File/Reg. Number</u>
32	Certification pursuant to Section 906 of the Sarbanes Oxley Act of 2002	X			
*	Management contract or compensatory plan or arrangement.				
**	Portions of this exhibit (indicated by asterisks) have been omitted pursuant to a request for confidential treatment and then filed separately with the SEC.				

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SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

AMEDICA CORPORATION

Date: March 31, 2014

/S/ Eric K. Olson

Eric K. Olson
Chief Executive Officer
(Principal Executive Officer)

Date: March 31, 2014

/S/ Jay M. Moyes

Jay M. Moyes
Chief Financial Officer
(Principal Financial and Accounting Officer)

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

Date: March 31, 2014

/S/ Eric K. Olson

Eric K. Olson, Director

Date: March 31, 2014

/S/ Jay M. Moyes

Jay M. Moyes, Director

Date: March 31, 2014

/S/ Max E. Link

Max E. Link, Ph.D. Director

Date: March 31, 2014

/S/ Sonny Bal

B. Sonny Bal, M.D., Director

Date: March 31, 2014

/S/ David W. Truetzel

David W. Truetzel, Director

Date: March 31, 2014

/S/ Jeffrey S. White

Jeffrey S. White, Director

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AMEDICA CORPORATION

INDEX TO CONSOLIDATED FINANCIAL STATEMENTS

Years ended December 31, 2013 and 2012

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Report of Independent Registered Public Accounting Firm

The Board of Directors and Stockholders of
Amedica Corporation

We have audited the accompanying consolidated balance sheets of Amedica Corporation as of December 31, 2013 and 2012, and the related consolidated statements of operations and comprehensive loss, statements of convertible preferred stock and stockholders' deficit, and cash flows for each of the two years in the period ended December 31, 2013. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

We conducted our audits in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. We were not engaged to perform an audit of the Company's internal control over financial reporting. Our audits included consideration of internal control over financial reporting as a basis for designing audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control over financial reporting. Accordingly, we express no such opinion. An audit also includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements, assessing the accounting principles used and significant estimates made by management, and evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the consolidated financial position of Amedica Corporation at December 31, 2013 and 2012, and the consolidated results of its operations and its cash flows for each of the two years in the period ended December 31, 2013, in conformity with U.S. generally accepted accounting principles.

The accompanying consolidated financial statements have been prepared assuming that the Company will continue as a going concern. As discussed in Note 1 to the financial statements, the Company has recurring losses from operations and needs to obtain additional financing in order to be compliant with debt covenants through 2014 that raise substantial doubt about its ability to continue as a going concern. Management's plans in regard to these matters are also described in Note 1. The consolidated financial statements do not include any adjustments that might result from the outcome of this uncertainty.

/s/ Ernst & Young LLP

Salt Lake City, Utah
March 31, 2014

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Amedica Corporation
Consolidated Balance Sheets
(in thousands, except share and per share data)

	December 31, 2013	December 31, 2012
Assets		
Current assets:		
Cash and cash equivalents	\$ 2,279	\$ 2,741
Restricted cash	392	260
Marketable securities	—	2,680
Trade accounts receivable, net of allowance of \$49 and \$58 respectively	2,817	4,016
Prepaid expenses and other current assets	1,575	519
Deferred offering costs	2,763	—
Inventories	10,084	8,826
Total current assets	19,910	19,042
Property and equipment, net	3,531	3,023
Intangible assets	4,688	5,189
Goodwill	6,163	6,163
Other long-term assets	35	38
Total assets	<u>\$ 34,327</u>	<u>\$ 33,455</u>
Liabilities and stockholders' deficit		
Current liabilities:		
Accounts payable	\$ 3,377	\$ 2,143
Accrued liabilities	3,711	1,606
Line of credit	—	2,573
Current portion of long-term debt	17,925	17,893
Total current liabilities	25,013	24,215
Deferred rent	575	606
Other long-term liabilities	134	134
Preferred stock warrant liability	11	526
Common stock warrant liability	199	2,783
Commitments and contingencies		
Convertible preferred stock, \$0.01 par value, 100,000,000 shares authorized; 80,910,394 and 76,170,394 shares issued and outstanding at December 31, 2013 and 2012; aggregate liquidation value of \$149,692 and \$140,212 at December 31, 2013 and 2012	161,456	153,474
Stockholders' deficit:		
Common stock, \$0.01 par value; 150,000,000 shares authorized; 597,675 and 348,636 shares issued and outstanding at December 31, 2013 and 2012	6	3
Additional paid-in capital / (capital deficiency)	(13,144)	(16,651)
Accumulated other comprehensive income	—	2
Accumulated deficit	(139,923)	(131,637)
Total stockholders' deficit	(153,061)	(148,283)
Total liabilities, convertible preferred stock and stockholders' deficit	<u>\$ 34,327</u>	<u>\$ 33,455</u>

See accompanying notes.

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Amedica Corporation
Consolidated Statements of Operations and Comprehensive Loss
(in thousands, except share and per share data)

	Year Ended December 31	
	2013	2012
Product revenue	\$ 22,314	\$ 23,065
Cost of revenue:		
Product revenue	5,752	5,423
Write-down of excess and obsolete inventory	1,293	1,043
Total cost of revenue	7,045	6,466
Operating expenses:		
Research and development	3,461	6,013
General and administrative	5,759	7,313
Sales and marketing	16,384	17,094
Impairment loss on intangible assets	—	15,281
Total operating expenses	25,604	45,701
Loss from operations	(10,335)	(29,102)
Other income (expense):		
Interest income	16	57
Interest expense	(1,851)	(5,611)
Loss on extinguishment of debt	—	(251)
Change in fair value of preferred stock warrants	515	(85)
Change in fair value of common stock warrants	3,455	(618)
Other expense	(86)	(151)
Total other income (expense)	2,049	(6,659)
Net loss before income taxes	(8,286)	(35,761)
Income tax benefit	—	726
Net Loss	(8,286)	(35,035)
Other comprehensive loss, net of tax:		
Unrealized gain (loss) on marketable securities	(2)	25
Total comprehensive loss	\$ (8,288)	\$ (35,010)
Net loss per share attributable to common stockholders:		
Basic and diluted	\$ (15.52)	\$ (100.52)
Weighted average common shares outstanding:		
Basic and diluted	534,073	348,550

See accompanying notes.

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Amedica Corporation
Consolidated Statements of Convertible Preferred Stock and Stockholders' Deficit
(in thousands, except share data)

	Convertible Preferred Stock		Common Stock		Additional Paid-In Capital/ (Capital Deficiency)	Accumulated Other Comprehensive Income (Loss)	Accumulated Deficit	Total Stockholders' Deficit
	Shares	Amount	Shares	Amount				
Balance at December 31, 2011	57,882,889	\$117,501	348,284	\$ 3	\$ (17,657)	\$ (23)	\$ (96,602)	\$ (114,279)
Issuance of Series C preferred stock as US Spine settlement shares	2,557,562	5,115	—	—	—	—	—	—
Issuance of Series E preferred stock as US Spine settlement shares	842,443	1,685	—	—	—	—	—	—
Issuance of Series F preferred stock upon conversion of convertible debt, net of issuance costs	14,887,500	29,173	—	—	—	—	—	—
Issuance of common stock upon exercise of stock options	—	—	450	—	5	—	—	5
Stock-based compensation	—	—	—	—	1,001	—	—	1,001
Unrealized gain on marketable securities	—	—	—	—	—	25	—	25
Net loss	—	—	—	—	—	—	(35,035)	(35,035)
Balance at December 31, 2012	76,170,394	\$153,474	348,734	\$ 3	\$ (16,651)	\$ 2	\$ (131,637)	\$ (148,283)
Issuance of Series F preferred stock for \$2.00 per share, net of issuance costs	4,740,000	7,982	—	—	—	—	—	—
Issuance of common stock in connection with exercise of warrants	—	—	178,516	2	2,876	—	—	2,878
Issuance of common stock upon cashless exercise of stock options	—	—	27,624	—	—	—	—	—
Issuance of common stock upon exercise of stock options	—	—	29,680	1	76	—	—	77
Stock-based compensation	—	—	13,121	—	555	—	—	555
Unrealized loss on marketable securities	—	—	—	—	—	(2)	—	(2)
Net loss	—	—	—	—	—	—	(8,286)	(8,286)
Balance at December 31, 2013	80,910,394	\$161,456	597,675	\$ 6	\$ (13,144)	\$ —	\$ (139,923)	\$ (153,061)

See accompanying notes.

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Amedica Corporation
Consolidated Statements of Cash Flows
(in thousands)

	Year Ended December 31	
	2013	2012
Cash flow from operating activities		
Net loss	\$ (8,286)	\$ (35,035)
Adjustments to reconcile net loss to net cash used in operating activities:		
Write-down of intangible assets	—	15,281
Depreciation expense	1,706	2,379
Amortization of intangible assets	501	1,885
Amortization of lease incentive for tenant improvements	20	20
Accretion of interest expense on US Spine-related note payable	—	142
Accretion of interest expense on new bank debt	431	4
Non-cash interest expense on convertible debt	—	2,512
Loss on extinguishment of debt	—	251
Non-cash interest expense on bank debt	—	22
Stock based compensation	555	1,001
Change in fair value of preferred stock warrant liability	(515)	85
Change in fair value of common stock warrant liability	(3,455)	618
Loss (gain) on sale of equipment	81	151
Write-down of excess and obsolete inventory	1,293	1,043
Bad debt expense	5	(28)
Changes in operating assets and liabilities:		
Trade accounts receivable	1,194	(541)
Prepaid expenses and other current assets	(4,220)	(844)
Inventories	(2,551)	1,529
Other long-term assets	3	—
Accounts payable and accrued liabilities	3,289	(200)
Deferred revenue	—	(5)
Net cash used in operating activities	(9,949)	(9,730)
Cash flows from investing activities		
Purchase of property and equipment	(2,332)	(593)
Proceeds from sale of property and equipment	37	—
Increase in restricted cash	(132)	(260)
Purchases of marketable securities	—	(5,082)
Proceeds from maturities of marketable securities	2,680	10,210
Net cash provided by investing activities	253	4,275
Cash flows from financing activities		
Proceeds from exercise of stock options and warrants	2,955	5
Proceeds from line of credit	10,940	2,573
Payments on line of credit	(13,513)	(2,000)
Proceeds from issuance of long-term debt	—	17,889
Issuance of preferred stock warrants	—	111
Payments on bank debt extinguishment	—	(15,500)
Payments on US Spine debtholder note	—	(3,000)
Proceeds from issuance of preferred stock	8,852	—
Proceeds from issuance of convertible debt and warrants, net	—	4,787
Net cash provided by financing activities	9,234	4,865
Net decrease in cash and cash equivalents	(462)	(590)
Cash and cash equivalents at beginning of year	2,741	3,331
Cash and cash equivalents at end of year	<u>\$ 2,279</u>	<u>\$ 2,741</u>
Supplemental cash flow information		
Common stock warrants issued in connection with preferred stock	\$ 871	\$ —
Cash paid for interest	\$ 1,403	\$ 3,079

See accompanying notes.

AMEDICA CORPORATION

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS
Years ended December 31, 2013 and 2012

1. Organization and Summary of Significant Accounting Policies

Amedica Corporation (“Amedica” or “the Company”) was incorporated in the state of Delaware on December 10, 1996. Amedica is a commercial-stage biomaterial company focused on using its silicon nitride technology platform to develop, manufacture, and commercialize a broad range of medical devices. The Company believes it is the first and only manufacturer to use silicon nitride in medical applications. The Company acquired US Spine, Inc. (“US Spine”), a Delaware spinal products corporation with operations in Florida, on September 20, 2010. The Company’s products are primarily sold in the U.S.

Basis of Presentation and Principles of Consolidation

These consolidated financial statements have been prepared by management in accordance with accounting principles generally accepted in the United States (“U.S. GAAP”), and include all assets and liabilities of the Company and its wholly-owned subsidiary, US Spine. All material intercompany transactions and balances have been eliminated.

Liquidity and Capital Resources

For the years ended December 31, 2013 and 2012, the Company incurred a net loss of \$8.3 million and \$35.0 million, respectively and used cash in operations of \$9.9 million and \$9.7 million, respectively. The Company had an accumulated deficit of \$139.9 million and \$131.6 million at December 31, 2013 and 2012, respectively. With the exception of a small net income for the years ended December 31, 2002 and 1999, the Company has incurred net losses in each year since inception. To date, the Company’s operations have been principally financed from proceeds from the issuance of preferred and common stock, convertible debt and bank debt and, to a lesser extent, cash generated from product sales. It is anticipated that the Company will continue to generate operating losses and use cash in operations through 2014.

As discussed further in Note 7, the Company is contractually obligated to repay \$18.0 million to a bank by making principal payments of \$600,000 over a period of 30 months beginning in January 2014. At December 31, 2013, the Company had approximately \$2.3 million in cash and investments. In order to finance growth in product sales, to invest in further product development and to otherwise satisfy obligations in 2014 as they mature, the Company needed to obtain additional financing. In February 2014, the Company completed an initial public offering (“IPO”), of its common stock, in which the Company sold and issued 3,682,900 shares, including 182,900 shares sold pursuant to the exercise by the underwriters of their over-allotment option, at an issuance price of \$5.75 per share, less underwriting discounts and commissions. As a result of the offering, the Company received proceeds of approximately \$15.4 million, net of approximately \$5.8 million in offering costs. The Company will need to obtain additional funding during the third quarter of 2014 to satisfy the \$7.2 million of principal obligations due under the GE Secured Lending Facility in 2014 (which amount excludes interest payments and any other fees incurred or that we may incur) and to maintain compliance with the financial and liquidity covenants related to the GE Secured Lending Facility through 2014. Furthermore, if the Company is unable to access additional funds prior to becoming non-compliant with the liquidity debt covenant, the entire remaining balance of the debt could become immediately due and payable at the option of the lender. The Company is seeking additional financing and expects to obtain additional financing through the issuance of common stock and/or debt, including a refinancing of its existing debt. Additional funding may not be available to the Company on acceptable terms, or at all. Any additional equity financing, if available to the Company, may not be available on favorable terms and will most likely be dilutive to its current stockholders, and debt financing, if available, may involve more restrictive covenants. The Company’s ability to access capital when needed is not assured and, if not achieved on a timely basis, will materially harm its business, financial condition and results of operations. These uncertainties create substantial doubt about the Company’s ability to continue as a going concern. No adjustment has been made to our financial statements as a result of this uncertainty. The Report of Independent Registered Public Accounting Firm at the beginning of these financial statements includes a going concern explanatory paragraph.

Reverse Stock Split

On February 11, 2014, the Company effected a 1 for 25.7746 reverse stock split of the Company’s common stock. The par value and the authorized shares of the common and convertible preferred stock were not adjusted as a result of the reverse stock split. All common stock share and per-share amounts for all periods presented in these financial statements have been adjusted retroactively to reflect the reverse stock split.

Use of Estimates

The preparation of financial statements in conformity with U.S. GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial

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statements and the reported amounts of revenue and expenses during the period. Actual results could differ from those estimates. Some of the more significant estimates relate to inventory, stock-based compensation, long-lived and intangible assets, contingent consideration and the liability for preferred stock and common stock warrants.

Concentrations of Credit Risk and Significant Customers

Financial instruments which potentially subject the Company to concentrations of credit risk consist primarily of cash and cash equivalents, marketable securities, accounts receivable and restricted cash. The Company limits its exposure to credit loss by placing its cash and cash equivalents with high credit-quality financial institutions in bank deposits, money market funds, U.S. government securities and other investment grade debt securities that have strong credit ratings. The Company has established guidelines relative to diversification of its cash and marketable securities and their maturities that are intended to secure safety and liquidity. These guidelines are periodically reviewed and modified to take advantage of trends in yields and interest rates and changes in the Company's operations and financial position. Although the Company may deposit its cash and cash equivalents with multiple financial institutions, its deposits, at times, may exceed federally insured limits.

The Company's customers are primarily hospitals and surgical centers. At December 31, 2013, one customer receivable balance was 15% of the Company's total trade accounts receivable. At December 31, 2012, no customer receivable balance was 10% or greater of the Company's total trade accounts receivable. There was one customer that accounted for 10% or more of the Company's revenue representing 14% of revenue for the years ended December 31, 2013 and 2012.

Credit to customers is granted based upon an analysis of the customers' individual credit worthiness. The Company's allowance for bad debts as of December 31, 2013 and 2012 was \$49,000 and \$58,000, respectively. For the years ended December 31, 2013 and 2012, the Company recorded bad debt expense (recovery) of \$5,000 and (\$28,000), respectively.

Revenue Recognition

The Company derives its product revenue primarily from the sale of spinal fusion devices and related products used in the treatment of spine disorders. The Company's product revenue is generated from sales to two types of customers: (1) surgeons and hospitals and (2) stocking distributors. Most of our products are sold on a consignment basis through a network of independent sales distributors; however, the Company also sells its products to independent stocking distributors. Product revenue is recognized when all four of the following criteria are met: (1) persuasive evidence that an arrangement exists; (2) delivery of the products has occurred; (3) the selling price of the product is fixed or determinable; and (4) collectability is reasonably assured. The Company generates the majority of its revenue from the sale of inventory that is consigned to independent sales distributors that sell the Company's products to surgeons and hospitals. For these products, the Company recognizes revenue at the time it is notified the product has been used or implanted and a valid purchase order has been received. For all other transactions, the Company recognizes revenue when title and risk of loss transfer to the stocking distributor, and all other revenue recognition criteria have been met. The Company generally recognizes revenue from sales to stocking distributors at the time the product is shipped to the distributor. Stocking distributors, who sell the products to their customers, take title to the products and assume all risks of ownership at time of shipment. The Company's stocking distributors are obligated to pay within specified terms regardless of when, if ever, they sell the products. The Company's policy is to classify shipping and handling costs billed to customers as an offset to total shipping expense in the statement of operations, primarily within sales and marketing. In general, the Company's customers do not have any rights of return or exchange.

Cost of Revenue

The expenses that are included in cost of revenue include all direct product costs and manufacturing costs. Specific provisions for excess or obsolete inventory are also included in cost of revenue. Beginning in January 2013, cost of revenue also includes the 2.3% excise tax on the sale of medical devices in the United States.

Cash, Cash Equivalents, Restricted Cash, and Investments

The Company considers all cash on deposit, money market accounts and highly-liquid debt instruments purchased with original maturities of three months or less to be cash and cash equivalents. Restricted cash consists of cash we receive from payments of our accounts receivables held in a segregated account that must be applied to pay amounts owed under our revolving credit facility with General Electric Capital Corporation. The Company's investments in marketable debt and equity securities are deemed by management to be available for sale and are reported at fair market value with the net unrealized appreciation or depreciation reported as a component of accumulated other comprehensive loss in stockholders' deficit. At the time of sale, any realized appreciation or depreciation, calculated by the specific identification method, is recognized in other income and expense.

Inventories

Inventories are stated at the lower of cost or market, with cost for manufactured inventory determined under the standard cost method which approximates first-in first-out ("FIFO"). Manufactured inventory consists of raw material, direct labor and

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manufacturing overhead cost components. Inventories purchased from third-party manufacturers are stated at the lower of cost or market using the first-in, first-out method. The Company reviews the carrying value of inventory on a periodic basis for excess or obsolete items, and records any write-down as a cost of revenue, as necessary. It is reasonably possible that the Company may be required to make adjustments to the carrying value of inventory in future periods. Inventory write-downs for excess or obsolete inventory are recorded as a cost of revenue. The Company holds consigned inventory at distributor and other customer locations where revenue recognition criteria have not yet been achieved.

Property and Equipment

Property and equipment, including surgical instruments and leasehold improvements, are stated at cost, less accumulated depreciation and amortization. Property and equipment are depreciated using the straight-line method over the estimated useful lives of the assets, which range from three to five years. Leasehold improvements are amortized over the shorter of their estimated useful lives or the related lease term, generally five years.

In accounting for long-lived assets, the Company makes estimates about the expected useful lives of the assets, the expected residual values of certain of these assets, and the potential for impairment based on the fair value of the assets and the cash flows they generate. The Company periodically evaluates the carrying value of long-lived assets to be held and used when events and circumstances indicate that the carrying amount of an asset may not be recovered. The Company has not recognized any impairment loss for property and equipment for the years ended December 31, 2013 and 2012.

Long-Lived Assets, Indefinite-lived Intangible and Goodwill

Periodically, the Company assesses potential impairment of its long-lived assets, which include property, equipment, and acquired intangible assets. The Company performs an impairment review whenever events or changes in circumstances indicate that the carrying value may not be recoverable. Factors the Company considers important which could trigger an impairment review include, but are not limited to, significant under-performance relative to historical or projected future operating results, significant changes in the manner of its use of acquired assets or its overall business strategy, and significant industry or economic trends. When the Company determines that the carrying value of a long-lived asset may not be recoverable based upon the existence of one or more of the above indicators, the Company determines the recoverability by comparing the carrying amount of the asset to net future undiscounted cash flows that the asset is expected to generate and recognizes an impairment charge equal to the amount by which the carrying amount exceeds the fair market value of the asset. The Company amortizes finite-lived intangible assets on a straight-line basis over their estimated useful lives. The Company recorded an impairment loss for definite-lived intangible assets of \$0 and \$13.4 million during the years ended December 31, 2013 and 2012, respectively.

The Company tests goodwill and indefinite-lived intangibles for impairment annually as of December 31, or whenever events or changes in circumstances indicate that goodwill or indefinitely-lived intangibles may be impaired. For goodwill, the Company initially assesses qualitative factors to determine whether the existence of events or circumstances leads to a determination that it is more-likely-than-not that the fair value of a reporting unit is less than its carrying amount. For goodwill impairment testing purposes, we consider the value of our equity, including the value of our convertible preferred stock, in the total carrying value of our single reporting unit. If, after assessing the totality of events or circumstances, the Company determines it is more-likely-than-not that the fair value of a reporting unit is less than its carrying amount, then the Company performs a first step by comparing the book value of net assets to the fair value of the Company's single reporting unit. If the fair value is determined to be less than the book value, a second step is performed to compute the amount of impairment as the difference between the estimated fair value of goodwill and the carrying value. The Company performed the first step and determined that it was not more-likely-than-not that the fair value of the Company's single reporting unit was less than its carrying amount and no goodwill impairment was recognized during the years ended December 31, 2013 or 2012.

The impairment test for indefinite-lived intangible assets not subject to amortization involves a comparison of the estimated fair value of the intangible asset, using the income approach, with its carrying value. If the carrying value of the intangible asset exceeds its fair value, an impairment loss is recognized in an amount equal to that excess. The Company recorded an impairment loss for indefinite-lived intangible assets of \$0 and \$1.9 million during the years ended December 31, 2013 and 2012, respectively.

Research and Development

All research and development costs, including those funded by third parties, are expensed as incurred. Research and development costs consist of engineering, product development, test-part manufacturing, testing, developing and validating the manufacturing process, and regulatory related costs. Research and development expenses also include employee compensation, employee and nonemployee stock-based compensation, supplies and materials, consultant services, and travel and facilities expenses related to research activities.

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Advertising Costs

Advertising costs are expensed as incurred. The primary component of the Company's advertising expenses is advertising in trade periodicals. Advertising costs were approximately \$403,000 and \$1.1 million for the years ended December 31, 2013 and 2012, respectively.

Income Taxes

The Company recognizes a liability or asset for the deferred tax consequences of all temporary differences between the tax basis of assets and liabilities and their reported amounts in the financial statements that will result in taxable or deductible amounts in future years when the reported amounts of the assets and liabilities are recovered or settled. The income tax benefit for the years ended December 31, 2013 and 2012 was \$0 and \$726,000, respectively. The Company recognizes interest and penalties as a component of the provision for income taxes. No interest or penalties were recognized in the years ended December 31, 2013 and 2012.

Stock-Based Compensation

The Company measures stock-based compensation expense related to employee stock-based awards based on the estimated fair value of the awards as determined on the date of grant and is recognized as expense over the remaining requisite service period. The Company utilizes the Black-Scholes-Merton option pricing model to estimate the fair value of employee stock options. The Black-Scholes-Merton model requires the input of highly subjective and complex assumptions, including the estimated fair value of the Company's common stock on the date of grant, the expected term of the stock option, and the expected volatility of the Company's common stock over the period equal to the expected term of the grant. The Company estimates forfeitures at the date of grant and revises the estimates, if necessary, in subsequent periods if actual forfeitures differ from those estimates.

The Company accounts for stock options and warrants to purchase shares of stock that are issued to non-employees based on the estimated fair value of such instruments using the Black-Scholes-Merton option pricing model. The measurement of stock-based compensation expense for these instruments is variable and subject to periodic adjustments to the estimated fair value until the awards vest or, in the case of convertible preferred stock warrants, each reporting period until the warrant is exercised. Any resulting change in the estimated fair value is recognized in the Company's consolidated statements of operations and comprehensive loss during the period in which the related services are rendered.

Deferred offering costs

Deferred offering costs totaled \$2.8 million at December 31, 2013. These costs represent legal and accounting costs related to the Company's efforts to raise capital through the IPO of the Company's common stock. There were no IPO costs incurred prior to May 1, 2013. Additional costs related to the Company's IPO activities were deferred until the completion of the IPO in February 2014, at which time they were reclassified to additional paid-in capital as a reduction of the IPO proceeds.

Net Loss Per Share

Basic net loss per share is calculated by dividing the net loss by the weighted-average number of common shares outstanding for the period, without consideration for common stock equivalents. Diluted net loss per share is calculated by dividing the net loss by the weighted-average number of common share equivalents outstanding for the period determined using the treasury-stock method. Dilutive common stock equivalents are comprised of convertible preferred stock, warrants for the purchase of convertible preferred stock and common stock, stock options and unvested RSUs outstanding under the Company's equity incentive plans. For all periods presented, there is no difference in the number of shares used to calculate basic and diluted shares outstanding due to the Company's net loss position.

Potentially dilutive securities not included in the calculation of diluted net loss per share because to do so would be anti-dilutive are as follows (in common stock equivalent shares):

	<u>Year ended December 31,</u>	
	<u>2013</u>	<u>2012</u>
Convertible preferred stock	3,884,788	3,578,513
Preferred stock warrants	102,810	90,971
Common stock options	106,544	287,418
Common stock warrants	473,835	377,578
Restricted stock units	168,832	—
Total	<u>4,736,809</u>	<u>4,334,480</u>

Shares are based on the terms of securities outstanding as of the date presented. If antidilution provisions are triggered for the Series F convertible preferred stock, additional dilution could occur.

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2. Intangible Assets

Indefinite lived intangible assets consist of trademarks, while amortizable intangible assets subject to amortization consist of customer relationships, developed technology and other patents and patent applications, all of which were acquired in 2010 in connection with the US Spine acquisition. The amortizable intangible assets are being amortized over a period of 12 years from the acquisition date.

Management of the Company noted that certain US Spine product sales, as well as sales to certain acquired US Spine customers during the one-year period ended December 31, 2012 were less than expected relative to the forecasted revenues at the time of the acquisition. This indicator prompted the Company to question whether the carrying value of its long-lived and indefinite lived intangible assets would be recoverable. Management compared the carrying amount of the assets to net future undiscounted cash flows that the intangible assets are expected to generate, and concluded that an impairment existed. All of the Company's definite-lived and indefinite-lived intangible assets are categorized within Level 3 of the fair value hierarchy. Management estimated the fair values of the intangible assets and recognized an impairment loss of approximately \$15.3 million in the year ended December 31, 2012. There was no impairment recorded in 2013. Significant assumptions used in valuing the intangibles in 2012 included the following:

Valuation technique:	Discounted cash flow method
Weighted-average cost of capital	17%
Weighted-average revenue growth rate	-58% to 10%
EBITDA margin	8.99% to 12.44%

The details of the impairment loss recorded in the year ended December 31, 2012 are presented below (in thousands):

Asset Group	Net Carrying Value of Intangible Assets Prior to Impairment	Impairment of Intangible Assets	Adjusted Net Carrying Value After Impairment Charge
Indefinite-Lived Intangible Assets			
Trademark - US SPINE	\$ 1,244	\$ (1,044)	\$ 200
Trademark - PREFERENCE	700	(550)	150
Trademark - FACET GUN/BOLT	205	(205)	—
Trademark - JAVELIN	100	(100)	—
Total indefinite-lived intangible assets	2,249	(1,899)	350
Long-lived intangible assets			
Customer Relationships	7,870	(5,780)	2,090
Developed technology	9,242	(6,788)	2,454
Other patents and patent applications	1,109	(814)	295
Total long-lived intangible assets	18,221	(13,382)	4,839
Total asset group	\$ 20,470	\$ (15,281)	\$ 5,189

Intangible assets consisted of the following at December 31, 2013 (in thousands):

	December 31, 2013		
	Carrying Amount	Accumulated Amortization	Intangible assets, net
Customer relationships	\$ 3,990	\$ 2,116	\$ 1,874
Developed technology	4,685	2,485	2,200
Other patents and patent applications	562	298	264
Trademarks	350	—	350
Total amortizable intangible assets	\$ 9,587	\$ 4,899	\$ 4,688

Based on the recorded intangibles at December 31, 2013, the estimated amortization expense for each of the five years ending in 2017 is approximately \$501,000 per year and \$1.8 million thereafter. The total accumulated amortization at December 31, 2012 was approximately \$4.4 million.

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3. Fair Value, Marketable Securities, and Contingent Consideration Liability

Fair Value Measurements

The Company has implemented the accounting requirements for financial assets, financial liabilities, non-financial assets and non-financial liabilities reported or disclosed at fair value. ASC 820, *Fair Value Measurements*, defines fair value, establishes a three level hierarchy for measuring fair value in generally accepted accounting principles, and expands disclosures about fair value measurements. Level 1 inputs are quoted prices (unadjusted) in active markets for identical assets or liabilities that a company has the ability to access at the measurement date. Level 2 inputs are inputs other than quoted prices included within Level 1 that are observable for the asset or liability, either directly or indirectly. Level 3 inputs are unobservable inputs for the asset or liability. This hierarchy requires the Company to use observable market data, when available, and to minimize the use of unobservable inputs when determining fair value. The methods described above may produce a fair value calculation that may not be indicative of net realizable value or reflective of future fair values. Furthermore, while the Company believes its valuation methods are appropriate and consistent with other market participants, the use of different methodologies or assumptions to determine the fair value of certain financial instruments could result in a different fair value measurement at the reporting date. Assets are classified in their entirety based on the lowest level of input that is significant to the fair value measurement.

The Company's financial instruments are cash and cash equivalents, marketable securities, accounts receivable, accounts payable, accrued liabilities, common and preferred stock warrant liabilities, and notes payable. The recorded values of cash and cash equivalents, accounts receivable, accounts payable, and accrued liabilities approximate their fair values based on their short-term nature. The fair value of marketable securities is primarily estimated based upon quoted market prices in either active or inactive markets. The fair value of the common stock warrant liability and the preferred stock warrant is estimated based upon a Black-Scholes-Merton option pricing model. The recorded value of notes payable approximates the fair value as the interest rate approximates market interest rates.

Marketable Securities

There were no marketable securities at December 31, 2013. Marketable securities at December 31, 2012 were as follows (in thousands):

	December 31, 2012			
	Amortized Cost	Gross Unrealized Gains	Gross Unrealized Losses	Estimated Fair Value
Certificates of deposit	\$ 1,000	\$ —	\$ —	\$ 1,000
Corporate debt securities	1,679	2	(1)	1,680
Total	\$ 2,679	\$ 2	\$ (1)	\$ 2,680

The majority of the Company's marketable securities were valued using quoted prices in markets that are not active or based on other observable inputs. Certificates of deposit and corporate debt securities are categorized as Level 2 because they are based on evaluated prices that reflect observable market information, such as actual trade information or similar securities, including interest rates and yield curves, adjusted for observable differences. The Company's asset manager obtains prices from pricing services, whose prices are obtained from direct feeds from market exchanges.

The Company had no marketable securities that were required to be re-measured at December 31, 2013. The following table sets forth the fair value of the Company's financial assets that were re-measured at December 31, 2012, based on the valuation approach applied to each class of security (in thousands):

	Level 1	Level 2	Level 3	Total
Certificates of deposit	\$ —	\$1,000	\$ —	\$1,000
Corporate debt securities	—	1,680	—	1,680
Total	\$ —	\$2,680	\$ —	\$2,680

There were no marketable securities at December 31, 2013.

No impairment losses were recognized through earnings related to available for sale securities for the years ended December 31, 2013 and 2012.

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The proceeds from maturities and sales of marketable securities and resulting realized gain and losses for the years ended December 31, 2013 and 2012 were as follows (in thousands):

	For the year ended December 31,	
	2013	2012
Proceeds from maturities and sales	\$ 2,680	\$ 10,210
Realized gains	2	—

For fair value disclosures regarding warrants to purchase convertible preferred stock, see Preferred Stock Warrant Liability under Note 8. For fair value disclosures regarding warrants to purchase common stock, see Common Stock Warrant Liability under Note 7.

4. Property and Equipment

The following is a summary of the components of property and equipment (in thousands):

	December 31,	
	2013	2012
Manufacturing and lab equipment	\$ 7,163	\$ 6,976
Surgical instruments	7,721	6,358
Leasehold improvements	1,439	1,430
Software and computer equipment	807	807
Furniture and equipment	621	621
	17,751	16,192
Less: accumulated depreciation and amortization	(14,220)	(13,169)
	<u>\$ 3,531</u>	<u>\$ 3,023</u>

5. Inventories

The following is a summary of the components of inventories (in thousands):

	December 31,	
	2013	2012
Raw materials	\$ 1,025	\$ 969
WIP	1,410	421
Finished Goods	7,649	7,436
	<u>\$10,084</u>	<u>\$8,826</u>

Finished goods include consigned inventory in the amounts of approximately \$5.5 million and \$5.6 million as of December 31, 2013 and 2012, respectively.

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6. Accrued Liabilities

Accrued liabilities consist of the following (in thousands):

	December 31,	
	2013	2012
Commissions	\$ 989	\$ 756
Payroll and related expenses	645	521
Royalties	277	246
Interest payable	119	72
Final loan payment fees	1,281	—
Offering costs	323	—
Deferred rent	32	7
Other	45	4
	<u>\$3,711</u>	<u>\$1,606</u>

7. Debt and Line of Credit

The following table summarizes the maturities of the Company's debt, at face value, at December 31, 2013 (in thousands):

2014	\$ 7,200
2015	7,200
2016	<u>3,600</u>
	<u>\$18,000</u>

Total interest expense for all debt instruments for the years ended December 31, 2013 and 2012 was \$2.4 million and \$5.6 million, respectively.

Bank Debt

In February 2009, the Company borrowed \$5.0 million from a bank under a promissory note agreement ("Term Loan 1"). The Term Loan agreement required that \$1.0 million in principal be repaid annually for five years, along with interest which was due and payable on a monthly basis. The interest rate was 3.5% per annum above the Ninety-Day London Interbank Offered Rate. In December 2009, the Company amended the agreement to extend the first principal repayment date to February 2010.

In April 2010, the Company further amended the Term Loan 1 agreement, extending the date of the first principal repayment, increasing the principal balance of the term loan note from \$5.0 million to \$7.5 million and adding a line of credit facility, with a total amount available of \$2.5 million. The Company received the proceeds from the additional \$2.5 million term loan in April 2010 and borrowed \$1.86 million under the line of credit in December 2010. In March 2011, the Company repaid the entire \$1.86 million borrowed under the line of credit. In connection with the amendment, the Company issued warrants to purchase 50,000 shares of the Company's Series E convertible preferred stock to the bank. The warrants became exercisable, after one year, at the purchase price of \$2.20 per share and expire after five years. There are certain conditions that would have caused the warrants to become immediately exercisable. In accordance with the provisions of ASC 480, *Distinguishing Liabilities from Equity*, the Company recorded a liability for the fair value of these warrants and records fair value adjustments to the liability at the end of each reporting period.

In September 2010, in connection with the acquisition of US Spine, the Company guaranteed a \$5.0 million loan to US Spine, a wholly owned subsidiary of the Company ("Term Loan 2"). Term Loan 2 matured in August 2015 and was secured by all of the assets of US Spine and required interest to be paid on a monthly basis.

In March 2011, the Company amended its Term Loan 1 and Term Loan 2 agreements. The March 2011 amendment to the Term Loan 1 agreement increased the principal from \$7.5 million to \$10.5 million, increased the annual payments from \$1.5 million to \$2.1 million and extended the date of the first principal repayment to April 1, 2012 (and annually thereafter) with a final payment due in April 2015. The March 2011 amendment to the Term Loan 2 extended the date of the first principal repayment of \$1.0 million to August 2012, with a final payment due in August 2015.

In May 2012, the Company amended its Term Loan 1 and Term Loan 2 agreements. The amendment did not change the principal balances, which remained at \$10.5 million and \$5.0 million, respectively; however, the annual principal repayments beginning in 2012 for both loans were waived and the maturity date was modified to be April 1, 2013. In connection with the May 2012 amendments new minimum gross profit and minimum EBITDA targets were established and the Company incurred approximately \$50,000 in legal and debt modification fees payable to the bank.

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In December 2012, the Company repaid all amounts outstanding under the Term Loan 1 and Term Loan 2 agreements with the bank, as well as all amounts outstanding under the line of credit facility, totaling \$18.0 million in principal and approximately \$36,000 in accrued interest. The Company wrote off approximately \$52,000 of capitalized debt issuance costs related to the term loans and the line of credit facility. The Company paid \$107,500 in commissions related to the debt restructuring, of which approximately \$70,000 was capitalized as debt issuance costs and the remaining \$37,500 was recorded as interest expense in 2012. See New Bank Debt disclosures below.

US Spine-Related Note Payable

In connection with the acquisition of US Spine in September 2010, the Company entered into a subordinated secured promissory note agreement to pay two equal payments of \$3.0 million to a US Spine stockholder over a two-year period. In accordance with this note agreement, the Company paid \$3.0 million to the US Spine stockholder during 2011 and paid the remaining \$3.0 million in 2012. The Company discounted the note using an imputed interest rate of 6.5%, which approximated a market rate of interest on the acquisition date.

Senior Secured Subordinated Convertible Promissory Notes

From March to May 2011, the Company issued \$24.8 million in aggregate principal amount of Senior Secured Subordinated Convertible Promissory Notes (the "Notes"). The Company also received a commitment to purchase \$5.0 million of Notes in February 2012, at the option of the Company ("Commitment Notes"). The Notes matured two years from issuance (one year on the Commitment Notes) and interest was paid quarterly at 6% for the first year and 8% for year two. The Notes may be converted to common stock at a conversion rate of \$51.55 per share. The Note holders also received 3-year warrants exercisable for shares of the Company's common stock equal to 50% of the principal amount of the Notes divided by two, with an exercise price of \$51.55 per share (total of 288,685 warrants) (the "Note Holder Warrants"). The warrants were immediately exercisable and expire after three years from issuance. The Company paid approximately \$1.2 million in commissions, loan fees and related costs in connection with the Notes, which were amortized over the two year term of the Notes. In connection with the closing of the Notes, and in accordance with the US Spine acquisition agreement, \$1.0 million of the \$24.8 million in proceeds was paid in March 2011 towards the \$3.0 million due to a US Spine stockholder in September 2011.

In December 2012, contemporaneous with the closing of the New Bank Debt (see below), holders of approximately 82% of the outstanding principal balance of the Notes consented to the conversion of the Notes into shares of Series F convertible preferred stock at \$2.00 per share, pursuant to the Company's request. All of the Notes were converted into 14,887,500 shares of Series F convertible preferred stock. See convertible preferred stock under Note 8.

New Bank Debt/Preferred Stock Warrant Liability

In December 2012, the Company closed on a \$21.5 million senior secured credit facility (the "New Bank Debt") with a bank consortium consisting of two lenders, one of which was the existing bank lender from whom the Company borrowed beginning in 2009 (see Bank Debt above). The new agreement consists of a term loan for \$18.0 million and a \$3.5 million revolving line of credit secured by the Company's accounts receivable, based on certain defined criteria, and matures in May 2016. The term loan consists of interest only payments for a period of 12 months after which monthly principal payments of approximately \$600,000 are required for a period of 30 months beginning in January 2014. The term loan bears interest at the fixed rate of 7.5% per annum, while the line of credit had an interest rate of 7.0% at December 31, 2013 and 2012, which is based on the variable rate of 5.5% plus the higher of (i) 1.5% and (ii) the three-month LIBOR determined as of two London business days divided by a number equal to 1.0 minus the aggregate of the rates of reserve requirements on the day that is two London business days prior to the beginning of the interest period for Eurocurrency funding that are required to be maintained by a member bank of the Federal Reserve System. There was no amount outstanding under the revolving line of credit and \$821,000 was available to borrow at December 31, 2013. The Company pledged all of its assets as collateral for the loan. The agreement includes a non-refundable final payment fee of \$720,000, as well as an annual management fee of \$15,000 per year.

The agreement includes certain financial covenants related to minimum liquidity including six (6) times the monthly cash burn amount, as defined, days sales outstanding of accounts receivable balances, and other financial reporting requirements. Upon any event of default, including the financial covenants, the lender may declare the loan immediately due and payable. The agreement provides for an unused credit facility fee of 0.75% per annum of the unused portion of the line of credit, payable monthly in arrears. The Company paid a total of approximately \$333,000 in fees and commissions associated with the New Bank Debt, of which approximately \$264,000 was capitalized as debt issuance costs and the remaining \$69,000 was recorded as interest expense in 2012. The agreement includes a final payment fee of \$720,000 due upon prepayment in full or at scheduled maturity.

In December 2012, in connection with the New Bank Debt, the Company issued warrants to purchase 270,000 shares of the Company's Series F convertible preferred stock to the two lenders. The warrants are immediately exercisable at the purchase price of \$2.00 per share and expire after ten years. The Company has accounted for these warrants under the provisions of ASC 480, *Distinguishing Liabilities from Equity*. Accordingly, the Company recorded a liability for the fair value of these warrants in 2012 and records fair value adjustments to the liability at the end of each reporting period. See Preferred Stock Warrant Liability under Note 8.

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Pursuant to its terms, the Company must repay the \$18.0 million New Bank Debt over a period of 30 months beginning in January 2014. As of December 31, 2013, the total outstanding principal was \$18.0 million, although the financial statements reflect a carrying value of \$17.9 million due to the bifurcated value of warrants issued in connection with the debt, which is being amortized to interest expense over the life of the loan. The Company had been in covenant default with regards to the liquidity covenant several times during 2013. The Company entered into four amendments to the agreement during 2013 to forego the liquidity covenant testing. The fourth amendment to the agreement entered into in December 2013 stipulated the liquidity covenant would not be tested through January 31, 2014. In January 2014, the Company entered into a fifth amendment to the agreement that extended the time through which the liquidity covenant would not be tested to February 28, 2014. Although the Company was in compliance with the liquidity covenant at February 28, 2014, the Company anticipates that it will be non-compliant with the liquidity covenant prior to December 31, 2014, and has therefore classified the entire obligation as a current liability. The Company incurred amendment fees which are being amortized to interest expense over the life of the loan. The total accrued amendment fees at December 31, 2013 were \$1.1 million. The Company expects to pay these amendment fees in April 2014. Except as noted above, the Company is in compliance with all other covenants under the agreement.

Common Stock Warrant Liability

The Company issued 288,685 Note Holder Warrants in 2011, which are considered mark-to-market liabilities which are re-measured to fair value at each reporting period due to a round down provision whereby the exercise price of the warrants would change, if subsequent equity instruments were issued with an exercise price less than \$51.55 per share. The Company also issued warrants to purchase 57,557 shares of common stock to the placement agent in connection with the issuance of the Notes in May 2011. These warrants are also considered mark-to-market liabilities which are re-measured to fair value at each reporting period.

The Company will continue to adjust the common stock warrant liability for changes in fair value until the earlier of the exercise of the warrants to purchase shares of common stock or the completion of a liquidation event. The common stock warrant liability fair value was determined using primarily unobservable inputs and has been classified as a Level 3 liability in accordance with U.S. GAAP.

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During 2012 the Company reduced the exercise price of the 288,685 Note Holder Warrants, from \$51.55 to \$25.77 per share, in connection with the conversion of the convertible debt into 14,887,500 shares of Series F convertible preferred stock as discussed above and in Note 8 and also extended the expiration date by four years to March 2018. This caused an increase in the value of the common stock liability of approximately \$2.0 million; see the table below. In February 2013, the Company further reduced the exercise price of the 288,685 Note Holder Warrants, from \$25.77 to \$17.53 per share, and offered a replacement warrant for every Note Holder Warrant exercised at \$17.53 per share. The replacement warrants contain the same terms and conditions (including exercise price and term) as before the February 2013 amendment. This caused an increase in the common stock liability of approximately \$339,000. As a result of this amendment, the Company raised \$3.1 million from the exercise of warrants for common stock and issued 178,516 shares of common stock. In connection with this financing, the Company paid Creation Capital LLC, its financial advisor, approximately \$250,000 in commissions pursuant to its 2012 financial advisor consulting agreement. During August and September 2013, the Company issued warrants to purchase a total of 101,262 shares of common stock in connection with the sale of Series F convertible preferred stock. See “Convertible Preferred Stock” under Note 8. These warrants are classified as liabilities which are re-measured to fair value at the conclusion of each reporting period due to a round down provision whereby the exercise price of the warrants would change if subsequent equity investments were issued with a purchase price less than \$25.77 per share with respect to 91,951 of the warrants and \$56.70 per share with respect to 9,311 of the warrants. The fair value of the warrants at the grant date was approximately \$870,000 based upon the Black-Scholes-Merton option pricing model and the assumptions set forth in the table below. This amount was recorded as a derivative liability and reduced the carrying amount of convertible preferred stock.

The following table summarizes the changes in the common stock warrant liability for the years ended December 31, 2013 and 2012 (in thousands):

	Common Stock Warrant Liability
Balance at December 31, 2011	\$ (2,165)
Increase in liability due to modification of warrant terms	(1,998)
Decrease in fair value included in earnings, as other income	1,380
Balance at December 31, 2012	(2,783)
Increase in liability due to issuance of warrants to placement agent	(55)
Increase in liability due to issuance of warrants to investors	(816)
Increase in liability due to modification of warrant terms	(424)
Decrease in fair value included in earnings, as other income	3,879
Balance at December 31, 2013	\$ (199)

The assumptions used in estimating the common stock warrant liability at December 31, 2013 and 2012 and for the years then ended, are set forth below:

	As of and for the year ended December 31,	
	2013	2012
Estimated fair value of Company common shares	\$5.75 to \$17.53	\$ 17.53
Weighted-average risk free interest rate	1.27%	0.68%
Weighted-average expected life (in years)	4.3	4.9
Expected dividend yield	0%	0%
Weighted average expected volatility	64%	71%

The \$5.75 issuance price of the common stock sold in the Company’s IPO in February 2014 was used as the estimated fair value of the Company’s common shares at December 31, 2013 due to the close proximity to the valuation date and there were no significant changes that would indicate a decrease or increase in the stock price. The price of \$17.53 was used as the estimated fair value of the Company’s common shares for estimated fair value calculations of the common stock warrant liability done as of and prior to September 30, 2013. The price of \$17.53 was based on a third party valuation performed as of September 30, 2012. The valuation technique used in the third party valuation was a hybrid of the discounted cash flow method and the guideline public company method. The significant assumptions used in determining the \$17.53 estimated fair value of the Company’s common shares were as follows:

Weighted-average cost of capital (WACC)	17%
Revenue growth rate (range)	32.5% to 5%
Compounded average revenue growth rate	17.7%
EBITDA margin (range)	(23.8)% to 32.7%

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The effect of changes to these significant assumptions on the estimated liability for common stock warrants are set forth below:

Fair value of Company common shares	Warrant liability increases
WACC increases	Warrant liability decreases
Revenue growth increases	Warrant liability increases
Average revenue growth increases	Warrant liability increases
EBITDA margin increases	Warrant liability increases
Risk free interest increases	Warrant liability decreases
Expected average life increases	Warrant liability increases
Expected dividend yield increases	Warrant liability decreases
Expected volatility increases	Warrant liability increases

8. Equity

Common Stock

The Company had reserved shares of common stock for future issuance as follows:

	December 31,	
	2013	2012
Convertible preferred stock:		
Shares outstanding	3,884,788	2,955,250
Warrants		
Series C convertible preferred stock	52,325	46,703
Series D convertible preferred stock	12,786	9,827
Series E convertible preferred stock	27,224	23,965
Series F convertible preferred stock	10,475	10,475
Common stock	473,835	377,578
Options outstanding	106,544	287,418
Restricted stock units	168,832	—
Shares available for grant under equity plans	91,659	160,124
	<u>4,828,468</u>	<u>3,871,340</u>

Shares are based on the terms of securities outstanding as of the date presented. Upon completion of the IPO in February 2014, antidilution provisions were triggered for the Series F convertible preferred stock and Series F warrants and the Series F preferred stock converted into 4,906,874 shares of common stock and the Series F warrants converted into warrants to purchase 67,499 shares of common stock. Upon completion of the IPO, all shares of preferred stock converted into 8,029,779 shares of common stock

Convertible Preferred Stock

In August and September 2013, the Company issued an aggregate of 4,740,000 shares of Series F convertible preferred stock at a purchase price of \$2.00 per share, and associated warrants to purchase 91,951 shares of common stock. The warrants have an exercise price of \$25.77 per share and expire after five years. In addition, the Company issued warrants to purchase an aggregate of 9,311 shares of common stock to a placement agent, in August and September 2013. These warrants contain an exercise price of \$56.70 per share and expire after five years. See “Common Stock Warrant Liability” under Note 7.

In December 2012, the Company issued 14,887,500 shares of Series F convertible preferred stock at \$2.00 per share upon conversion of the Notes. The placement agent received commissions of approximately \$447,000 in connection with this conversion. See Senior Secured Subordinated Convertible Promissory Notes under Note 7. See also “Conversion Price Protection for Series F Convertible Preferred Stock” below.

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At December 31, 2013, convertible preferred stock consisted of the following:

<u>Series</u>	<u>Designated Shares</u>	<u>Shares Issued and Outstanding</u>	<u>Aggregate Liquidation Preference (in thousands)</u>
Series A	5,800,000	5,365,398	\$ 3,219
Series A-1	10,400,000	10,390,463	6,234
Series B	2,300,000	1,944,147	2,333
Series B-1	3,300,000	3,299,141	3,959
Series C	7,900,000	6,682,562	13,365
Series C-1	4,325,000	4,275,000	8,550
Series D	8,300,000	7,978,800	23,936
Series D-1	6,300,000	6,145,667	18,437
Series E	16,200,000	15,201,716	30,404
Series F	20,710,000	19,627,500	39,255
Undesignated	14,465,000	—	—
Total	<u>100,000,000</u>	<u>80,910,394</u>	<u>\$ 149,692</u>

At December 31, 2012, convertible preferred stock consisted of the following:

<u>Series</u>	<u>Designated Shares</u>	<u>Shares Issued and Outstanding</u>	<u>Aggregate Liquidation Preference (in thousands)</u>
Series A	5,800,000	5,365,398	\$ 3,219
Series A-1	10,400,000	10,390,463	6,234
Series B	2,300,000	1,944,147	2,333
Series B-1	3,300,000	3,299,141	3,959
Series C	7,900,000	6,682,562	13,365
Series C-1	4,325,000	4,275,000	8,550
Series D	8,300,000	7,978,800	23,936
Series D-1	6,300,000	6,145,667	18,437
Series E	16,200,000	15,201,716	30,404
Series F	14,950,000	14,887,500	29,775
Undesignated	20,225,000	—	—
Total	<u>100,000,000</u>	<u>76,170,394</u>	<u>\$ 140,212</u>

The rights and preferences of the convertible preferred stock are as follows:

Dividends

Holders of the convertible preferred stock shall be entitled to receive noncumulative dividends in preference to any dividend on common stock payable only if declared by the Board of Directors. As of December 31, 2013 and 2012, the Board of Directors had not declared any dividends.

Liquidation Preference

In the event of any liquidation or winding up of the Company, including in the event of the merger, consolidation and sale of the Company, the holders of Series F convertible preferred stock shall be entitled to receive, in preference to holders of all other series of preferred stock and holders of common stock, a per share amount equal to \$2.00, plus all accrued but unpaid dividends, when, as and if declared. If the Series F convertible preferred stock liquidation preference is paid in full to holders of such preferred stock, thereafter, the holders of Series A and A-1, Series B and B-1, Series C and C-1, Series D and D-1 convertible preferred stock, and Series E convertible preferred stock, shall be entitled to receive ratably, and in preference to the holders of common stock, a per share amount equal to \$0.60 for Series A and A-1 convertible preferred stock, \$1.20 for Series B and B-1 convertible preferred stock, \$2.00 for Series C and C-1 convertible preferred stock, \$3.00 for Series D and D-1 convertible preferred stock, and \$2.00 for Series E convertible preferred stock plus, with respect to each such series of preferred stock, all declared but unpaid dividends. After the payment of the liquidation preference to the holders of the preferred stock, the remaining assets shall be distributed ratably to the holders of the common stock.

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A sale, merger, reorganization, liquidation, dissolution or winding up of the Company may, in certain circumstances, be deemed to be a liquidation event and trigger the liquidation preferences associated with the outstanding shares of convertible preferred stock. Because a change in control could occur and not be solely within the control of the Company, all convertible preferred stock has been deemed to be redeemable and classified outside of permanent equity in the accompanying consolidated balance sheets. However, because the timing of any such redemption is uncertain, the Company will not accrete the carrying value of the convertible preferred stock to its liquidation preference value until it becomes probable that redemption will occur.

Conversion

The holders of the convertible preferred stock shall have the right to convert the shares of preferred stock held by such holders, at any time, into shares of common stock. Upon conversion, any declared but unpaid dividends on the preferred stock will be paid in additional shares of common stock.

The convertible preferred stock shall be automatically converted into common stock, at the then applicable conversion ratio, upon the closing of a public offering of shares of common stock at a per share price not less than the then applicable conversion price (as adjusted for stock splits, stock dividends, recapitalizations, etc.).

The conversion ratio of each series of convertible preferred stock at December 31, 2013, was as noted below:

<u>Series</u>	<u>Conversion Ratio</u>
Series A	0.0388
Series A-1	0.0582
Series B	0.0414
Series B-1	0.0591
Series C	0.0435
Series C-1	0.0631
Series D	0.0505
Series D-1	0.0653
Series E	0.0441
Series F	0.0388

Conversion Price Protection for Series F Convertible Preferred Stock

Holders of the Series F convertible preferred stock received certain protections in the event the Company closes an IPO or closes a transaction that constitutes a change in control of the Company. Upon completion of the IPO in February 2014, all 19,627,500 shares of Series F convertible preferred stock converted into 4,906,874 shares of common stock.

Voting Rights

The preferred stock will vote together with the common stock, and not as separate classes, except as specifically provided below or as otherwise required by law. Each share of preferred stock shall have a number of votes equal to the number of shares of common stock the preferred stock is convertible into on an as-converted basis.

Unless an affirmative vote of 50% of the combined outstanding shares of preferred stock, voting separately as a class, is obtained, the Company shall not undertake any of the following: (i) declaration or payment of any dividend or other distribution or payment on the (or the redemption, purchase or other acquisition for value of any) capital stock of the Company or any subsidiary; (ii) any liquidation, dissolution, recapitalization or reorganization of the Company; (iii) transfer or disposition of assets or rights with a value of more than \$1,000,000; and/or (iv) any amendment to the Company's certificate of incorporation that changes or alters any of the preferences, voting powers or other rights and privileges of preferred stock.

Registration Rights

The preferred stockholders and warrant holders were granted registration rights that provide these holders the right to request, beginning 180 days after the completion of a qualifying initial public offering, that the Company file a registration statement to register under the Securities Act the common stock that would be issued upon conversion of the preferred shares or exercise of the warrants. Thereupon, the Company is obligated to use commercially reasonable efforts to timely file a registration statement. These registration rights are subject to certain conditions and limitations, including the Company's right, based on advice of the lead managing underwriter of a future offering, to limit the number of shares included in any such registration under certain circumstances.

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Preferred Stock Warrant Liability

In connection with the various convertible preferred stock offerings, the placement agent received warrants to purchase convertible preferred stock. The warrants are fully exercisable after one year from issuance and expire after seven years. As described in Note 11, Related-Party Transactions, during 2012 the Company extended the expiration date for the warrants to purchase Series C convertible preferred stock by an additional five years. The exercise price of these warrants is equal to 110% of the offering price of the underlying convertible preferred stock. On the closing of the initial public offering, these warrants will convert into warrants to purchase shares of common stock at the then applicable conversion rate for the related preferred stock. During 2012, the Company granted warrants to purchase Series F convertible preferred stock at \$2.00 per share to a bank in connection with the debt refinancing (see Note 7, New Bank Debt). The warrants to purchase Series F convertible preferred stock are immediately exercisable and expire after ten years. The grant dates, number of warrants, exercise price and estimated fair value of the warrants at December 31, 2013 and 2012, were as noted below:

<u>Series of convertible preferred stock underlying warrants</u>	<u>Grant Date</u>	<u>Number of Warrants</u>	<u>Exercise Price</u>	<u>Estimated Fair Value at December 31, 2013 (in thousands)</u>
Series C	02/24/06	1,203,750	\$ 2.20	\$ —
Series D	04/27/07	253,290	\$ 3.30	—
Series E	03/22/10	617,691	\$ 2.20	—
Series F	12/17/12	270,000	\$ 2.00	11
		<u>2,344,731</u>		<u>\$ 11</u>

<u>Series of convertible preferred stock underlying warrants</u>	<u>Grant Date</u>	<u>Number of Warrants</u>	<u>Exercise Price</u>	<u>Estimated Fair Value at December 31, 2012 (in thousands)</u>
Series C	02/24/06	1,203,750	\$ 2.20	\$ 277
Series D	04/27/07	253,290	\$ 3.30	8
Series E	03/22/10	617,691	\$ 2.20	129
Series F	12/17/12	270,000	\$ 2.00	111
		<u>2,344,731</u>		<u>\$ 525</u>

Freestanding warrants for shares that are either puttable or warrants for shares that are redeemable are classified as liabilities on the balance sheet at fair value. In connection with the grant of the warrants to purchase Series C convertible preferred stock in 2006, Series D convertible preferred stock in 2007, Series E convertible preferred stock in 2010, and Series F convertible preferred stock in 2012, the Company recorded the initial fair values of the warrants of approximately \$929,000, \$442,000, \$266,000, and \$111,000, respectively, as a preferred stock warrant liability. At the end of each reporting period, changes in fair value during the period are recorded as a component of other income or expense. The preferred stock warrant liability fair value was determined using primarily unobservable inputs and has been classified as a Level 3 liability.

For the years ended December 31, 2013 and 2012, the Company recorded a benefit of \$514,000 and a charge of \$85,000, respectively, for the change in fair value of the preferred stock warrants. The Company will continue to adjust the liability for changes in fair value until the earlier of the exercise of the warrants to purchase shares of convertible preferred stock, the completion of a liquidation event, including the completion of the initial public offering, at which time the liabilities will be reclassified to stockholders' deficit when the warrants are converted to common stock warrants, or the expiration of the warrants.

The valuation of the preferred stock warrant liability has been determined using the underlying common share value; See Common Stock Warrant Liability under Note 7. The following table presents the changes in the estimated fair value of the preferred stock warrant liability for the years ended December 31, 2013 and 2012 (in thousands):

	<u>Preferred Stock Warrant Liability</u>
Balance at December 31, 2011	\$ (329)
Increase in liability due to issuance of warrants	(112)
Change in fair value included in earnings	(85)
Balance at December 31, 2012	(526)
Change in fair value included in earnings	515
Balance at December 31, 2013	<u>\$ (11)</u>

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The assumptions used in estimating the preferred stock warrant liability at December 31, 2013 and 2012 and for the years then ended, are set forth below:

	As of and for the year ended December 31,	
	2013	2012
Estimated fair value of Company common shares	\$5.75 to \$17.53	\$ 17.53
Weighted-average risk free interest rate	1.11%	0.68%
Weighted-average expected life (in years)	3.6	4.9
Expected dividend yield	0%	0%
Weighted average expected volatility	44%	71%

The \$5.75 issuance price of the common stock sold in the Company's IPO in February 2014 was used as the estimated fair value of the Company's common shares at December 31, 2013 due to the close proximity to the valuation date and there were no significant changes that would indicate a decrease or increase in the stock price. The price of \$17.53 was used as the estimated fair value of the Company's common shares for estimated fair value calculations of the common stock warrant liability done as of and prior to September 30, 2013. The price of \$17.53 was based on a third party valuation performed as of September 30, 2012. The valuation technique used in the third party valuation was a hybrid of the discounted cash flow method and the guideline public company method. The significant assumptions used in determining the \$17.53 estimated fair value of the Company's common shares were as follows:

Weighted-average cost of capital (WACC)	17%
Revenue growth rate (range)	32.5% to 5%
Compounded average revenue growth rate	17.7%
EBITDA margin (range)	(23.8)% to 32.7%

The effect of changes to these significant assumptions on the estimated liability for preferred stock warrants are set forth below:

Fair value of Company common shares	Warrant liability increases
WACC increases	Warrant liability decreases
Revenue growth increases	Warrant liability increases
Average revenue growth increases	Warrant liability increases
EBITDA margin increases	Warrant liability increases
Risk free interest increases	Warrant liability decreases
Expected average life increases	Warrant liability increases
Expected dividend yield increases	Warrant liability decreases
Expected volatility increases	Warrant liability increases

2003 and 2012 Option and Equity Plans

Under the Company's 2003 Stock Option Plan (the "2003 Plan"), the Company's Board of Directors has authorized the grant of options to employees and nonemployees for the issuance of up to 465,575 shares of the Company's common stock. All options granted have a term of ten years from the date of the grant and generally become fully exercisable within four years of continued employment or service at a rate defined in each option agreement.

In September 2012, the Company's Board of Directors adopted the 2012 Employee, Director and Consultant Equity Incentive Plan (the "2012 Plan") and resolved to cease awarding any further equity awards under the 2003 Plan. At that time there were approximately 155,192 approved and available shares of Common Stock available for issuance under the 2003 Plan. The Board resolved to transfer the available approximately 155,192 shares under the 2003 Plan into the 2012 Plan and further resolved that any outstanding shares of Common Stock represented by awards previously granted under the 2003 Plan that are forfeited, expire or are cancelled without delivery of shares of Common Stock shall be made available for award under the 2012 Plan.

The total number of shares available for grant under the 2012 Plan at December 31, 2013 was 91,659.

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Stock Options

A summary of the Company's stock option activity for the year ended December 31, 2013 was as follows:

	Options	Weighted-Average Exercise Price	Weighted-Average Remaining Contractual Life (Years)	Intrinsic Value
Outstanding at December 31, 2012	287,417	\$ 23.20		
Granted	13,191	\$ 25.77		
Exercised	(60,718)	\$ 2.71		
Cancelled	(133,346)	\$ 24.47		
Outstanding at December 31, 2013	106,544	\$ 28.90	4.7	\$ —
Exercisable at December 31, 2013	101,185	\$ 29.07	4.7	\$ —
Vested and expected to vest at December 31, 2013	106,419	\$ 28.90	4.9	\$ —

The aggregate intrinsic value in the table above is calculated as the difference between the estimated fair value of the Company's stock at December 31, 2013 and the exercise price of each option.

The weighted average grant date fair value of options granted during the years ended December 31, 2013 and 2012 was \$0.07 and \$14.18, respectively. The total intrinsic value of options exercised during the years ended December 31, 2013 and 2012 was approximately \$900,000 and \$6,400, respectively, calculated as the difference between the exercise price of the underlying stock option awards and the estimated fair value of the Company's common stock on the date of exercise. Cash received from option exercises for the years ended December 31, 2013 and 2012 was approximately \$77,000 and \$5,300, respectively. The Company recorded no tax benefit related to options exercised during the years ended December 31, 2013 and 2012.

The Company estimates the fair value of each stock option on the grant date using the Black-Scholes-Merton valuation model, which requires several estimates including an estimate of the fair value of the underlying common stock. The board of directors estimated the fair value of the Company's common stock on grant dates, taking into consideration a number of factors including a third-party valuation of the common stock, the issuance price of shares of the Company's convertible preferred stock, the preferential terms and conditions of the convertible preferred stock, the status of scientific research and development efforts and associated milestones and the likelihood of achieving a liquidity event for the share of the Company's common stock. The expected volatility was based on an average of the historical volatility of a peer group of similar companies. The expected term was calculated utilizing the simplified method. The risk-free interest rate was based on the U.S. Treasury yield curve in effect at the time of grant for the expected term of the option. No options were granted to employees during the year ended December 31, 2013. The following weighted average assumptions were used in the calculation to estimate the fair value of options granted to employees during the year ended December 31, 2012:

Weighted-average risk-free interest rate	1.14%
Weighted-average expected life (in years)	5.34
Expected dividend yield	0%
Weighted-average expected volatility	72%
Weighted-average fair value of the underlying stock	\$17.53

In March 2012, the board of directors of the Company approved the cancellation of stock options held by active employees and board members with exercise prices above \$25.77 per share, and the replacement of such options with options for the same number of shares with 100% immediate vesting on the date of grant. As a result, 47,100 options were modified resulting in recording additional stock-based compensation expense of approximately \$413,000 during the year ended December 31, 2013. The following table summarizes the Black-Scholes-Merton valuation inputs used in calculating the fair value and incremental expense recognized:

	Inputs Before Modification	Inputs Following Modification
Weighted-average risk-free interest rate	0.50%	1.11%
Weighted-average expected life (in years)	2.5	5.0
Expected dividend yield	0%	0%
Weighted-average expected volatility	65%	71%

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Restricted Stock Units

Restricted stock unit (“RSU”) activity for the year ended December 31, 2013 was as follows:

	<u>Number of Awards</u>	<u>Weighted-Average Grant Date Fair Value</u>
Unvested at December 31, 2012	—	
Granted	197,335	\$ 13.65
Vested	(19,399)	\$ 5.67
Forfeited	(9,104)	\$ 17.53
Unvested at December 31, 2013	<u>168,832</u>	<u>\$ 14.36</u>

The total fair value of RSUs vested during the year ended December 31, 2013 was \$110,000. No RSUs vested during the years ended December 31, 2012.

In February 2013, employees of the Company elected to exchange 93,968 stock options for an equal number of RSUs pursuant to a one-time tender offer approved by the board of directors. The RSUs vest solely upon either a change in control or upon the expiration of a lock-up period in connection with an underwritten public offering of shares of the Company’s common stock. The incremental fair value on the date of the exchange between the original stock options and the RSUs issued of approximately \$758,000 will be recognized when vesting conditions are achieved.

Stock-Based Awards Granted to Nonemployees

The Company issued 13,121 shares of common stock to nonemployees for sales and marketing services during the year ended December 31, 2013, and recorded \$231,000 in stock-based compensation expense during the year ended December 31, 2013.

The Company from time to time grants options to purchase common stock to non-employees for services. The Company estimates the fair value of these stock options using the Black-Scholes-Merton valuation model at each reporting date and records expense ratably over the vesting period of each stock option award. No options were granted to non-employees during the year ended December 31, 2012. The Company granted 13,191 options to non-employees and recorded stock-based compensation expense of \$17,000 during the year ended December 31, 2013.

The following assumptions were used in the Black-Scholes-Merton valuation model related to non-employee stock options granted during the year ended December 31, 2013:

Weighted-average risk-free interest rate	2.74%
Weighted-average expected life (in years)	10.0
Expected dividend yield	0%
Weighted-average expected volatility	52%
Weighted-average fair value of the underlying stock	\$5.75

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Summary of Stock-Based Compensation Expense

Total stock-based compensation expense included in the consolidated statements of comprehensive income was allocated as follows (in thousands):

	Years Ended December 31,	
	2013	2012
Cost of revenue	\$ —	\$ 31
Research and development	56	158
General and administrative	224	480
Selling and marketing	273	188
Capitalized into inventory	2	144
Total stock-based compensation expense	<u>\$ 555</u>	<u>\$ 1,001</u>

Unrecognized stock-based compensation for stock options and RSUs at December 31, 2013 was as follows (in thousands):

	Unrecognized Stock-Based Compensation	Weighted Average Remaining Period of Recognition (in years)
Stock options	\$ 336	1.6
RSUs	1,666	0.6
Total unrecognized stock-based compensation at December 31, 2013	<u>\$ 2,002</u>	

9. Income Taxes

The income tax benefit consisted of the following (in thousands):

	Years Ended December 31,	
	2013	2012
Current:		
Total current	\$ —	\$ —
Deferred:		
State	—	(81)
Federal	—	(645)
Total deferred	—	(726)
Total benefit	<u>\$ —</u>	<u>\$ (726)</u>

Deferred income taxes reflect the net tax effects of temporary differences between the carrying amount of assets and liabilities for financial reporting purposes and the amounts used for income tax purposes.

The following is a reconciliation of the expected statutory federal income tax provision to the actual income tax benefit (in thousands):

	Years Ended December 31,	
	2013	2012
Federal statutory rate	(34.0)%	(34.0)%
State taxes, net of federal benefit	(4.3)%	(4.3)%
Research and development credits	(3.0)%	(0.7)%
Equity related expenses	(5.0)%	6.7%
Change in valuation allowance	46.3%	30.3%
Total income tax expense	<u>0.0%</u>	<u>(2.0)%</u>

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Significant components of the Company's deferred tax assets were as follows (in thousands):

	December 31,	
	2013	2012
Deferred tax assets:		
Net operating loss carryforwards	\$ 42,661	\$ 38,610
Depreciation	17	90
Research credits	2,350	2,113
Other	2,823	3,329
Total deferred tax assets	47,851	44,142
Deferred tax liabilities:		
Amortization of intangible assets	(1,028)	(1,138)
Total deferred tax liabilities	(1,028)	(1,138)
Net deferred tax assets	46,823	43,004
Less valuation allowance	(46,957)	(43,138)
Net deferred tax assets (liabilities)	\$ (134)	\$ (134)

At December 31, 2013 and 2012, the Company had net operating loss carryforwards for federal and state income tax purposes of approximately \$111.5 million and \$99.1 million, respectively. The federal and state net operating loss carryforwards will expire from 2023 to 2033, unless previously utilized.

The income tax benefit of \$726,000 recorded in 2012 relates to the impairment of intangible assets associated with the 2010 US Spine acquisition. In accordance with Section 382 of the Internal Revenue Code, a change in ownership of greater than 50% within a three-year period places an annual limitation on the Company's ability to utilize its existing net operating loss carryforwards. The Company may be subject to these annual limitations and, therefore, unable to fully utilize the net operating loss carryforwards. Additionally, the Company may be unable to fully utilize all of the net operating loss carryforwards associated with the US Spine acquisition, due to certain annual limitations.

During the years ended December 31, 2013 and 2012, the Company recognized no amounts related to tax interest or penalties related to uncertain tax positions. The Company is subject to taxation in the United States and various state jurisdictions. The Company currently has no years under examination by any jurisdiction.

A valuation allowance has been established as realization of such deferred tax assets has not met the more likely-than-not threshold requirement. If the Company's judgment changes and it is determined that the Company will be able to realize these deferred tax assets, the tax benefits relating to any reversal of the valuation allowance on deferred tax assets will be accounted for as a reduction to income tax expense. The tax valuation allowance increased by approximately \$3.8 million and \$11.3 million for the years ended December 31, 2013 and 2012, respectively.

10. Commitments and Contingencies

The Company currently leases laboratory, manufacturing and office space and equipment under noncancelable operating leases which provide for rent holidays and escalating payments. Lease incentives, including rent holidays, allowances for tenant improvements and rent escalation provisions, are recorded as deferred rent. Rent under operating leases is recognized on a straight-line basis beginning with lease commencement through the end of the lease term. For each of the years ended December 31, 2013 and 2012, rental expense was approximately \$830,000 and \$810,000, respectively.

The following table summarizes future minimum rental payments required under operating leases that have initial or remaining non-cancelable lease terms in excess of one year as of December 31, 2013 (in thousands):

Years ending December 31:	
2014	\$ 856
2015	883
2016	910
2017	938
2018	962
Thereafter	980
Total minimum lease payments	<u>\$5,529</u>

The Company has entered into consulting and development agreements with some of its advisors, including some surgeon advisors. The Company has agreed to pay some of the surgeon advisors a portion of the net profits attributable to the sale of specific spine

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products for which the surgeon advisors provided the Company with consulting and related services related to the conceptualization, development, testing, clearance, approval and/or related matters involving implant products. The Company is obligated to pay royalties to as many as 14 different surgeon advisors in connection with the sale of certain of its implant products. These agreements generally continue until the later of (a) ten years from the date of the agreements, and (b) the expiration of the patent rights relating to the devices covered by the agreements, when rights have been assigned by the individuals to the Company. The Company incurred royalties of \$892,000 and \$852,000 related to these agreements for the years ended December 31, 2013 and 2012, respectively. None of the royalty arrangements contain minimum royalty payments.

The Company has executed agreements with certain executive officers of the Company which, upon the occurrence of certain events related to a change in control, call for payments to the executives up to three times their annual salary and accelerated vesting of previously granted stock options.

From time to time, the Company is subject to various claims and legal proceedings covering matters that arise in the ordinary course of its business activities. Management believes any liability that may ultimately result from the resolution of these matters will not have a material adverse effect on the Company's consolidated financial position, operating results, or cash flows.

11. Related-Party Transactions

Gregg R. Honigblum, the Chief Executive Officer of each of Creation Capital, LLC ("Creation Capital") and Creation Capital Advisors, LLC ("Creation Advisors"), joined the Company's board of directors in December 2006 and resigned in September 2013. The Company completed offerings of preferred stock and convertible debt through Creation Capital, as its placement agent, and it received strategic financial advisory services from Creation Advisors.

In February 2012, the Company received the final tranche of \$5.0 million in Notes (see Note 8) and paid Creation Capital as the Company's placement agent.

In connection with the conversion of the Notes to shares of Series F convertible preferred stock in December 2012, the Company was obligated to pay Creation Advisors a strategic financial advisory fee of approximately \$447,000. Creation Advisors agreed to a payment plan whereby the Company would pay one half of the advisory fee (or approximately \$224,000) immediately, and pay the other half in monthly installments over a period of 24 months. Also pursuant to the financial advisor consulting agreement, the Company entered into with Creation Advisors in June 2012, the Company paid Creation Advisors a strategic financial advisory fee of approximately \$108,000 in 2012 related to the new bank financing which closed in December 2012 (see Note 7).

Pursuant to the June 2012 financial advisor consulting agreement with Creation Advisors, the Company agreed to extend the expiration date of all warrants to purchase Series C convertible preferred stock previously issued to Creation Capital from February 2013 to February 2018. In connection with this modification, the Company recorded additional expense of approximately \$520,000 during the year ended December 31, 2012.

In conjunction with the warrant restructuring and the sale and issuance of other shares of common stock in March 2013, Creation Advisors was paid a strategic financial advisory fee of approximately \$250,000 (see Note 7—Common Stock Warrant Liability). In October 2013, the Company entered into a one-year consulting agreement for financial advisory services with Creation Advisors in which Creation Advisors will receive compensation of up to \$180,000 in cash (payable \$15,000 per month). The Company paid \$45,000 during the year ended December 31, 2013 under this agreement.

12. 401(k) Plan

Effective June 1, 2004, the Company adopted a defined contribution retirement plan under Section 401(k) of the Internal Revenue Code. The plan covers substantially all employees. Eligible employees may contribute amounts to the plan, via payroll withholdings, subject to certain limitations. The plan permits, but does not require, additional matching contributions to the plan by the Company on behalf of the participants in the plan. The Company incurred approximately \$162,000 and \$151,000 relating to retirement contributions for the years ended December 31, 2013 and 2012, respectively.

13. Subsequent Events

In November 2013 the Company was in default with respect to the minimum liquidity covenant under its agreement with a bank. The Company obtained an amendment to the agreement in December 2013 which stipulated that the liquidity covenant would not be tested through January 31, 2014. The Company obtained an additional amendment to the agreement on January 28, 2014 from the bank that expires on March 1, 2014. This amendment extended the time through which the liquidity covenant would not be tested to February 28, 2014.

In January 2014, the Board increased the aggregate number of shares issuable under the 2012 Plan to 3,000,000. In January 2014, the Board approved grants under the 2012 Plan totaling 1,692,980 RSUs to be issued on effectiveness of the filing of a registration statement on Form S-8 and vest upon the expiration of the lock-up period for our IPO. The estimated aggregate value to be recognized as compensation expense in 2014 for the RSUs granted on January 27, 2014 is \$9.7 million.

On February 12, 2014, the Company completed an IPO of its common stock, in which the Company sold and issued 3,682,900 shares, including 182,900 shares sold pursuant to the exercise by the underwriters of their over-allotment option, at an issuance price of \$5.75 per share, less underwriting discounts and commissions. As a result of the offering, the Company received proceeds of approximately \$15.4 million, net of approximately \$5.8 million in underwriting and offering costs.

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On February 11, 2014, the holders of a majority of the outstanding shares of the Company's Series F convertible preferred stock agreed to waive the conversion adjustment under the Company's Restated Certificate of Incorporation such that in no event will the denominator used to calculate the conversion ratio be less than \$8.00, provided that the Company completes its IPO on or before June 30, 2014. Upon completion of the IPO in February 2014, all 80,910,394 shares of preferred stock converted into 8,029,779 shares of common stock. Furthermore, all outstanding warrants exercisable for 2,344,731 shares of preferred stock converted into warrants exercisable for 159,834 shares of common stock.

The table below shows, on a pro forma basis, the impact of our IPO on certain consolidated balance sheet items. The pro forma condensed consolidated balance sheet data below gives effect to (i) the sale of 3,682,900 shares of common stock from the IPO at an offering price of \$5.75 per share after deducting the underwriting discounts and commissions and offering expenses incurred by the Company, (ii) conversion of all the outstanding shares of convertible preferred stock into 8,029,779 shares of common stock, and (iii) conversion of the preferred stock warrants into warrants to purchase a total of 159,834 shares of common stock and the related reclassification of the convertible preferred stock warrant liability to stockholders' equity upon the completion of the offering (in thousands):

	December 31, 2013	Pro Forma December 31, 2013 (unaudited)
Condensed consolidated balance sheet data:		
Cash and cash equivalents	\$ 2,279	\$ 17,648
Total current assets	19,910	35,279
Total assets	34,327	49,696
Preferred stock warrant liability	11	—
Convertible preferred stock	161,456	—
Common stock	6	123
Additional paid-in capital (deficiency)	(13,144)	163,575
Total stockholders' equity (deficit)	(153,061)	23,775
Total liabilities, convertible preferred stock and stockholders' equity (deficit)	\$ 34,327	\$ 49,696

CERTIFICATION OF CHIEF FINANCIAL OFFICER

I, Jay M. Moyes, certify that:

1. I have reviewed this annual report on Form 10-K of Amedica Corporation;

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 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)) for the registrant and we 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) 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

(c) 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 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 function):

(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 controls over financial reporting.

Date: March 31, 2014

/s/ Jay M. Moyes

Jay M. Moyes
Chief Financial Officer

CERTIFICATIONS UNDER SECTION 906

Pursuant to section 906 of the Sarbanes-Oxley Act of 2002 (subsections (a) and (b) of section 1350, chapter 63 of title 18, United States Code), each of the undersigned officers of Amedica Corporation., a Delaware corporation (the "Company"), does hereby certify, to such officer's knowledge, that:

The Annual Report for the year ended December 31, 2013 (the "Form 10-K") of the Company fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, and the information contained in the Form 10-K fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: March 31, 2014

/s/ Eric K. Olson

Eric K. Olson
Chief Executive Officer

Date: March 31, 2014

/s/ Jay M. Moyes

Jay M. Moyes
Chief Financial Officer