

# 2021 Annual Report



**Greg Rockingham**  
Jazz Drummer

Inspire Day:  
May 10, 2019

On the cover:

# Greg Rockingham



## Greg's Inspire Story

When Greg got Inspire in 2019, he had been living with sleep apnea for two decades. If it wasn't sleep apnea ruining Greg's sleep, it was his schedule. He's a jazz drummer, and a renowned one at that. His band Soul Message Band plays weekly at Chicago's legendary Green Mill Cocktail Lounge. His previous band, Deep Blue Organ Trio, opened for Steely Dan for many years. On his own, Greg has kept the beat for the likes of Charles Earland and Ellis Marsalis. Now, with Inspire, Greg's quality of life has improved, and he's at the top of his drumming game. "I have to play, and I have to have Inspire to get some rest. That's about the only two things I have to do in life."

No mask. No hose. Just sleep.™



# A Letter from the CEO

2021 Annual Report  
Inspire Medical Systems, Inc.

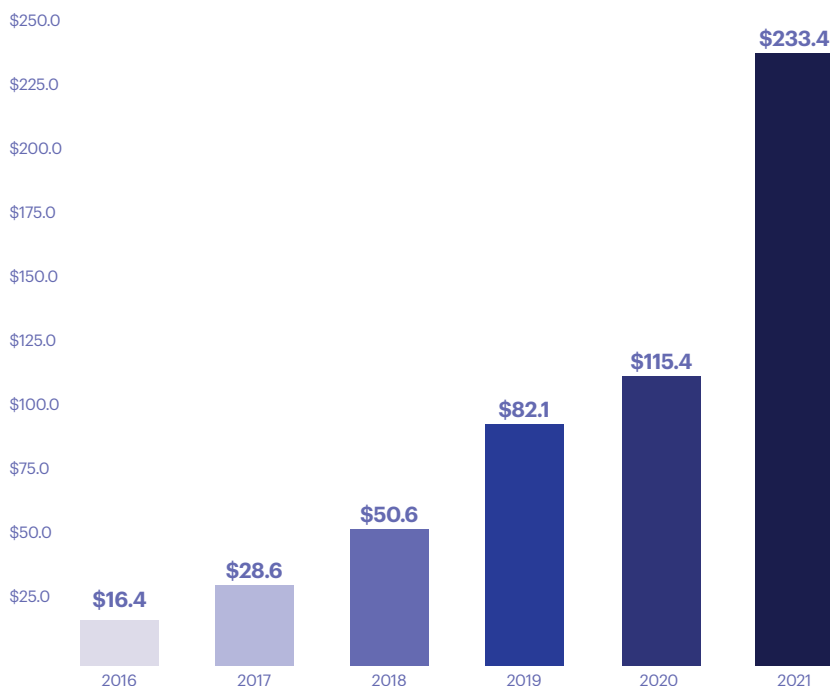
## Dear Fellow Stockholders,

In 2021, the continuing COVID-19 pandemic required significant operational flexibility globally by the Inspire team and healthcare providers in order to provide patients with Inspire therapy. The global lockdowns experienced in 2020 evolved into a series of local surges in infections last year and, as such, we were better positioned to anticipate and prepare for these events. Our purposeful approach minimized the impact that the pandemic had on Inspire procedures during 2021 and resulted in extremely strong operating performance, with full-year revenue of \$233.4 million, which represented growth of 102% over the \$115.4 million generated in 2020. We also increased our gross margin to 85.7% in 2021, as compared to 84.7% in 2020, and 83.4% in 2019.



**Timothy P. Herbert**  
President & Chief Executive Officer

### Revenue (in millions)



**\$233.4M**  
revenue  
in 2021

**102%**  
revenue  
growth  
in 2021

Since our FDA approval in 2014, we have continued to execute on our commercial strategy that is focused primarily on expanding Inspire's U.S. footprint, while also delivering strong growth in select key European countries. In the U.S., we strive to balance our growth between increasing utilization of our therapy at existing implanting centers, as well as training and activating new centers, with the ultimate objective of generating added capacity, thereby improving access for patients.

Historically, about half of our growth has been the result of increasing the number of patients treated at existing centers, and the remainder was generated by training and activating new centers. In 2020, the COVID-19

(continued on page 2)

**85.7%**  
gross  
margin  
in 2021



**259 new  
U.S.  
centers**

initiated Inspire programs in 2021

**684 U.S.  
centers**

in total at the end of 2021

pandemic required the shutdown of all centers for varying durations throughout the first half of the year. This limited the growth that we may have otherwise seen from existing centers compared to 2019. As such, in 2020, our growth was predominantly the result of opening new centers. In 2021, we experienced more balanced growth, with existing centers contributing approximately 60% of our growth. The team will continue to focus on driving balanced growth in the adoption of Inspire therapy, increasing utilization at existing implanting centers while also activating new centers.

In 2021, we trained and activated 259 new U.S. centers, ending the year with 684 U.S. implanting centers, up over 60% year-over-year. To reiterate, increasing the number of new centers will remain a priority going forward. With the improved reimbursement environment and because it is an outpatient procedure, ambulatory surgical centers (ASCs) are highly attractive settings where Inspire procedures can occur. Therefore, we expect that the number of ASCs offering Inspire therapy will continue to increase at a slightly higher rate as compared to hospitals. To meet this demand, Inspire has, and will continue to, enter into national pricing agreements with select national and regional ASC networks, which significantly reduces the logistical burden of opening new sites. Of course, we intend to maintain our focus on activating new centers in both hospitals and ASCs.

Another key element in increasing patient access to Inspire therapy is linked to our ability to assist prospective patients with making a connection to a qualified healthcare provider through our InspireSleep.com website and our advisor care program (ACP). The ACP is a call center that can provide potential patients with information about Inspire therapy, available commercial and Medicare insurance coverage, and make a tailored referral to an ENT surgeon or sleep physician in the patient's local area.

The ACP was created in 2020 and had an immediate positive impact on patient access and awareness of Inspire therapy. Anticipating increased demand for the ACP, we transitioned to a new larger vendor in early Q3 of this past year to enhance the technology features of the program and expand it to additional centers. By the end of 2021, over 550 implanting centers, or 80% of our total centers, were accessible through the ACP, up from just 180 the previous year. We are extremely pleased with the program and intend to continue expanding our ACP throughout 2022 and will implement additional technology enhancements intended to improve our ability to help patients connect with qualified healthcare providers, including through use of the Inspire Sleep app and the Inspire Cloud patient management system.

In order to succeed in our focus areas, we need to ensure that we have adequate support in the field, as well as a training team equipped to add additional centers, all while maintaining strong patient outcomes, which is always our top priority. The rate at which we open new sales territories is directly proportional to the rate at which we open new centers. As such, in 2021 we opened 50 new U.S. territories, ending the year with 157 U.S. territories. We also increased the field support teams, including augmenting the number of field clinical representatives, who specialize in case coverage and programming training.



In preparation for our planned continued U.S. expansion, we restructured our sales management leadership team by increasing the number of Area Vice Presidents (AVPs) to eight from four at the end of 2020. Each of these AVPs will be partnered with and manage an Area Business Manager (ABM), whose primary responsibility is to recruit and open new implanting centers. By doubling the number of ABMs, we significantly increased our ability to continue opening new centers at our desired growth rate.

In 2021, our market access team continued to work closely with the Center for Medicare & Medicaid Services (CMS) and the American Academy of Otolaryngology (AAO), which is the ENT physician society, to establish appropriate payment levels for the new billing codes for Inspire therapy, which became effective on January 1, 2022. As background, in November of 2020, a new category I code for the entire Inspire implant was approved by the American Medical Association. Further, a new category I code was also approved for the drug-induced sleep endoscopy procedure, which is used by physicians to assess whether a patient is a candidate for Inspire therapy. The permanent reimbursement levels for physicians, hospitals and ASCs have now been implemented, thereby removing another significant barrier for physicians and centers to initiate Inspire therapy programs.

Moreover, during 2021, we saw continued momentum in the expansion of positive coverage decisions. We ended 2021 with approximately 260 million lives under policy in the U.S. The majority of commercial payors now have positive coverage for Inspire therapy, as well as Medicare coverage in all 50 states and at Veteran Affairs (VA) and military hospitals. With this success in establishing coverage and reimbursement for Inspire therapy, we are well-positioned to further scale our commercial activity in the U.S.

In addition to our emphasis on the U.S. market, we are pleased with the recent growth in the adoption of Inspire therapy in Europe and other key markets. Our strategy is to focus our commercial activities in those countries that have established reimbursement, primarily Germany and the Netherlands. We experienced positive changes in both of these countries in 2021. Most notably, in Germany, we received permanent reimbursement with the DRG, the German hospital reimbursement system, which became effective on January 1, 2021. We will also continue to pursue reimbursement in other key European countries and are focused on growing activity in the United Kingdom, France, and Belgium, as well as expanding into other parts of Europe.

Following receipt of regulatory approval in Japan, we worked diligently in 2021 to establish appropriate reimbursement levels in that country and entered into an exclusive distribution agreement with Japan Lifeline, a leading medical technology distributor in Japan. Despite significant impact from the COVID pandemic, our internal training team was able to travel to Japan and support the first Inspire therapy procedure in that country in early 2022. We look forward to further expansion in Japan, as well as potentially leveraging expansion opportunities in select countries throughout the Asia Pacific region.

(continued on page 4)

80%  
of our  
centers

were accessible through the ACP  
by the end of 2021

50  
new U.S.  
territories

opened in 2021

157 U.S.  
territories

in total at the end of 2021



# Over 190 peer- reviewed papers

documenting the safety and  
efficacy of Inspire therapy

# New FDA- approved patient remote

launches in 2022, which is  
Bluetooth® enabled



The Inspire team continues to be fully committed to delivering positive and consistent outcomes for patients with untreated Obstructive Sleep Apnea (OSA). Monitoring and reporting on these outcomes provides an opportunity for continued education of ENT surgeons and sleep physicians, which, in turn, can lead to enhanced patient outcomes. To this end, in 2021, independent and company-supported researchers authored over 40 peer-reviewed publications on Inspire therapy, increasing the total number of such publications to over 190. These articles were published in both major ENT surgical and sleep medicine journals.

Looking ahead, we have a strong balance sheet to support our growth objectives. As of December 31, 2021, our cash, cash equivalents and investments were \$224.4 million. As such, we remain focused on expanding the capacity of our sales organization to meet the growing demand for Inspire therapy and advancing a number of key R&D projects.

Specifically, we are continuing with the development of the Inspire V next generation neurostimulation system and the next phase of the Inspire digital patient data management system, the Inspire Cloud. We are targeting the submission of the Inspire V PMA supplement application in late 2022. In 2022, we will introduce the new FDA-approved patient remote, which is Bluetooth® enabled and facilitates the transmission of patient and device information to the Inspire Cloud via the patient's smartphone and the Inspire Sleep app. We are targeting full body MRI approval by mid-2022 and also intend to complete development of the next-generation physician programmer this year.

In closing, while 2021 was another challenging year due to the impact of the pandemic, our team remained committed to helping patients and clinicians, and showed tremendous resolve during this tumultuous period, which resulted in a very successful year that saw us surpass 20,000 patients receiving Inspire therapy. On behalf of the entire Inspire team, we look forward to helping more people discover the potentially life-changing impact that Inspire therapy can offer patients with OSA and their loved ones. We sincerely thank all of our investors for their support and remain firmly committed to creating additional long-term stockholder value.

Sincerely,

A handwritten signature in blue ink, appearing to read 'Timothy P. Herbert'.

Timothy P. Herbert  
President & Chief Executive Officer

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

FORM 10-K

(Mark One)

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

For the fiscal year ended December 31, 2021

or

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

For the transition period from to

Commission File Number: 001-38468



**Inspire Medical Systems, Inc.**

(Exact name of registrant as specified in its charter)

**Delaware**

(State or other jurisdiction of incorporation or organization)

**26-1377674**

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

**5500 Wayzata Blvd., Suite 1600  
Golden Valley, MN**

(Address of principal executive offices)

**55416**

(Zip Code)

**(844) 672-4357**

(Registrant's telephone number, including area code)

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

Title of each class	Trading Symbol(s)	Name of exchange on which registered
<b>Common stock, \$0.001 par value</b>	<b>INSP</b>	<b>New York Stock Exchange</b>

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

**None**

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

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or 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 every Interactive Data File required to be submitted 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 such files). Yes ☒ No ☐



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

Large accelerated  
filer ☒

Accelerated  
filer ☐

Non-accelerated  
filer ☐

Smaller reporting  
company ☐

Emerging growth  
company ☐

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

Indicate by check mark whether the registrant has filed a report on and attestation to its management's assessment of the effectiveness of its internal control over financial reporting under Section 404(b) of the Sarbanes-Oxley Act (15 U.S.C. 7262(b)) by the registered public accounting firm that prepared or issued its audit report. ☒

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

As of June 30, 2021, the last business day of the registrant's most recently completed second fiscal quarter, the aggregate market value of the registrant's common stock held by non-affiliates was approximately \$5,161,346,505, based on the closing price of the registrant's common stock as reported on the New York Stock Exchange on such date.

As of February 7, 2022, the registrant had 27,529,438 shares of common stock, \$0.001 par value per share, outstanding.

#### **DOCUMENTS INCORPORATED BY REFERENCE**

Portions of the registrant's definitive proxy statement for its 2022 annual stockholders' meeting, which is to be filed within 120 days of the registrant's fiscal year ended December 31, 2021, are incorporated by reference into Part III of this Annual Report on Form 10-K.

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## 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. We intend such forward-looking statements to be covered by the safe harbor provisions for forward-looking statements contained in 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 facts contained in this Annual Report on Form 10-K are forward-looking statements, including statements regarding our future results of operations and financial position, business strategy, the impact of the ongoing and global COVID-19 pandemic on our business, financial results and financial position, prospective products, international product approvals and commercializations, our expectations regarding the final reimbursement levels for Inspire therapy procedures, research and development costs, timing and likelihood of success, other insurance providers' plans to begin approving our Inspire therapy, and the plans and objectives of management for future operations.

In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "expect," "plan," "anticipate," "could," "intend," "target," "project," "contemplate," "believe," "estimate," "predict," "potential" or "continue" or the negative of these terms or other similar expressions, although not all forward-looking statements contain these words. The forward-looking statements in this Annual Report on Form 10-K are only predictions and are based largely on our current expectations and projections about future events and financial trends that we believe may affect our business, financial condition, and results of operations. These forward-looking statements speak only as of the date of this Annual Report on Form 10-K and are subject to a number of known and unknown risks, uncertainties and assumptions, including, but not limited to the factors described in "Part I, Item 1. Business," "Part I, Item 1A. Risk Factors," and "Part I, Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations" of this Annual Report on Form 10-K.

Moreover, we operate in an evolving environment. New risk factors and uncertainties may emerge from time to time, and it is not possible for management to predict all risk factors and uncertainties.

Except as required by applicable law, we do not plan to publicly update or revise any forward-looking statements contained herein, whether as a result of any new information, future events, changed circumstances or otherwise.

Unless the context requires otherwise, references to "Inspire," the "Company," "we," "us," and "our," refer to Inspire Medical Systems, Inc.

## Summary Risk Factors

Our business is subject to numerous risks and uncertainties, including those described in Part I, Item 1A. "Risk Factors" in this Annual Report on Form 10-K. You should carefully consider these risks and uncertainties when investing in our common stock. The principal risks and uncertainties affecting our business include the following:

- our history of operating losses and dependency on our Inspire system for revenues;
- commercial success and market acceptance of our Inspire therapy;
- our ability to achieve and maintain adequate levels of coverage or reimbursement for our Inspire system or any future products we may seek to commercialize;
- competitive companies and technologies in our industry;
- the impact on our business, financial condition, and results of operation from the ongoing and global COVID-19 pandemic, or any other pandemic, epidemic or outbreak of an infectious disease;
- our ability to expand our indications and develop and commercialize additional products and enhancements to our Inspire system;



- future results of operations, financial position, research and development costs, capital requirements, and our needs for additional financing;
- our ability to forecast customer demand for our Inspire system and manage our inventory;
- our dependence on third-party suppliers and contract manufacturers;
- risks related to consolidation in the healthcare industry;
- our ability to expand, manage, and maintain our direct sales and marketing organization, and to market and sell our Inspire system in markets outside of the United States;
- our ability to manage our growth;
- our ability to hire and retain our senior management and other highly qualified personnel;
- risks related to product liability claims and warranty claims;
- our ability to address quality issues that may arise with our Inspire system;
- our ability to successfully integrate any acquired business, products or technologies;
- changes in global macroeconomic conditions;
- any failure of key information technology systems, processes or sites or damage to or inability to access our physical facilities;
- our ability to commercialize or obtain regulatory approvals or certifications for our Inspire therapy and system, or the effect of delays in commercializing or obtaining regulatory approvals or certifications;
- any violations of anti-bribery, anti-corruption, and anti-money laundering laws;
- risks related to our indebtedness;
- our ability to use our net operating losses and research and development carryforwards;
- the risk that we may be deemed to be an investment company under the Investment Company Act of 1940;
- U.S. Food and Drug Administration ("FDA") or other United States or foreign regulatory actions affecting us or the healthcare industry generally, including healthcare reform measures in the United States and international markets;
- our ability to establish and maintain intellectual property protection for our Inspire therapy and system or avoid claims of infringement; and
- risks related to our common stock.

## PART I

### Item 1. Business.

#### Overview

We are a medical technology company focused on the development and commercialization of innovative, minimally invasive solutions for patients with obstructive sleep apnea ("OSA"). Our proprietary Inspire system is the first and only FDA-approved neurostimulation technology that provides a safe and effective treatment for moderate to severe OSA. We have developed a novel, closed-loop solution that continuously monitors a patient's breathing and delivers mild hypoglossal nerve stimulation to maintain an open airway. A significant body of clinical data, which includes a publication in the *New England Journal of Medicine* and more than 190 peer-reviewed publications, supports the safety and efficacy of Inspire therapy. Inspire therapy received premarket approval ("PMA") from the FDA in April 2014 and has been commercially available in certain European markets since November 2011. Japan's Ministry of Health, Labour and Welfare ("MLHW") approved Inspire therapy to treat moderate to severe OSA in June 2018 and was formally added to the Japan National Health Insurance Payment Listing in June 2021. Inspire therapy is indicated for patients with moderate to severe OSA who do not have significant central sleep apnea and do not have a complete concentric collapse of the airway at the soft palate level. In addition, patients in the United States ("U.S.") and Japan must have been confirmed to fail or be unable to tolerate positive airway pressure ("PAP") treatments, such as continuous positive airway pressure ("CPAP"), and be 18 years of age or older, though there are no similar requirements for patients in Europe. Physicians have treated more than 20,000 patients with Inspire therapy at over 765 medical centers across the U.S., Europe, and Japan.

Sleep apnea is a serious and chronic disease that negatively impacts a patient's sleep, health, and quality of life. OSA is the most common form of sleep apnea. OSA occurs when a person's breathing is interrupted during sleep by a partially or completely blocked airway and affects patients of all ages, sexes, and body types. The severity of OSA is measured by the number of partial or complete airway blockages that a patient experiences in an hour, referred to as the apnea-hypopnea index ("AHI"). Moderate OSA patients have an AHI of 15 to 30 events per hour, while severe OSA patients have an AHI of 30 or more events per hour. Left untreated, OSA increases the risk of high blood pressure, hypertension, heart failure, stroke, coronary artery disease, and other life-threatening diseases.

CPAP is the leading therapy for patients with moderate to severe OSA. CPAP is delivered through a face or nasal mask that connects through a hose to a bedside air pump. In order for CPAP to be most effective, the mask must form an airtight seal on the patient's face or nose and the mask must be worn every night. The effectiveness of CPAP has been limited by low patient compliance as many patients find the mask or treatment cumbersome, uncomfortable, and loud. When CPAP fails or cannot be tolerated, patients' remaining treatment options consist primarily of invasive surgical procedures developed to modify or remove existing tissue in an attempt to create free air flow. These invasive surgical procedures have limited or unpredictable clinical benefit, are irreversible, and can be extremely painful. We believe that there is both an urgent clinical need and a strong market opportunity for an alternative to CPAP that is effective and minimally invasive.

Inspire therapy is an innovative, closed-loop, minimally invasive solution that provides comfort and convenience, resulting in high compliance for patients with moderate to severe OSA. Once implanted, the Inspire system delivers electrical stimulation that causes a slight forward movement of the back of the tongue, which helps to maintain an open airway, enabling the patient to inhale freely without interruption. We believe our Inspire therapy provides the following benefits:

- **Safe, effective, and durable treatment** supported by compelling clinical data, including long-term efficacy results out to five years from initial treatment.
- **Closed-loop system** that uses a proprietary algorithm to continuously monitor patients' breathing and provide electrical stimulation during the inspiratory phase.

- **Comfortable and convenient therapy resulting in high patient satisfaction** that was reported to be 91% at an average of 12 months from initial treatment in the first 1,849 patients in our ongoing global patient registry.
- **Strong patient compliance**, with 80% of patients reporting continued nightly use through five years from initial treatment in our Stimulation Therapy for Apnea Reduction ("STAR") trial.
- **Minimally invasive outpatient procedure** with short recovery time.
- **Long-lasting solution** with a battery designed to last approximately 11 years without charging or maintenance.

The results from multiple clinical trials, which include seven sponsored and more than 70 independent clinical studies that evaluated several thousand patients, have shown that our Inspire therapy provides statistically significant and sustained reduction in the severity of patients' OSA, improvement in sleep-related quality of life and reduction in snoring, as well as high patient compliance rates and a strong safety profile.

Our pivotal STAR trial was designed to demonstrate longitudinal therapy efficacy and included a randomized controlled therapy withdrawal study. The longitudinal study demonstrated an approximately 70% reduction in the median AHI in patients with moderate to severe OSA from a baseline of 29.3 events per hour to 9.0 events per hour at 12 months following initial treatment. STAR trial follow-up has shown results similar to the initial data at 18 months, three years, and five years. At five years, median AHI in patients with moderate to severe OSA remained low at 6.2 events per hour. The effectiveness of Inspire therapy was further demonstrated by the results of the randomized controlled therapy withdrawal study, in which patients in the therapy withdrawal group regressed to near-baseline AHI levels while patients in the control group that continued therapy experienced sustained therapeutic benefits.

We sell our Inspire system to hospitals and ambulatory surgery centers ("ASCs") in the U.S. and in select countries in Europe through a direct sales organization, and we sell our Inspire system in Japan through a distributor. As of December 31, 2021, we had 157 sales territories in the U.S. and 12 in Europe. Our direct sales force engages in sales efforts and promotional activities focused on ear, nose and throat ("ENT") physicians, and sleep centers. In addition, we highlight our compelling clinical data and value proposition to increase awareness and adoption amongst referring physicians. We build upon this top-down approach with strong direct-to-consumer marketing initiatives to create awareness of the benefits of our Inspire system and drive demand through patient empowerment. This outreach helps to educate thousands of patients on our Inspire therapy.

Our customers are generally reimbursed for the cost of patient treatment by various third-party payors, such as commercial insurance providers and Medicare. As of February 15, 2022, we have secured positive coverage policies with most U.S. commercial payors, including virtually all large national commercial insurers, encompassing approximately 260 million covered lives in the U.S. We are in active discussions with regional commercial insurers to establish additional positive coverage policies, as well as modify existing positive coverage policies to support reimbursement of Inspire therapy. In parallel, a subset of our 20-person reimbursement team, which we refer to as our market access team, is focused on assisting patients and physicians in obtaining prior authorization approvals from commercial payors on a case-by-case basis in advance of treatment with our Inspire therapy. In addition, all seven Medicare Administrative Contractors ("MACs") published final policies in 2020 that provide coverage of Inspire therapy when certain coverage criteria are met. We also have a U.S. government contract for patients who are treated by the Veterans Health Administration.

The procedure performed to implant our device was previously described for billing purposes using a Category I Current Procedural Terminology ("CPT") code (64568), which was used in conjunction with a temporary Category III CPT code (+0466T). At the October 2020 American Medical Association ("AMA") CPT Editorial Panel meeting, the AMA approved the creation of new Category I CPT codes (64582, 64583, and 64584) to separately identify hypoglossal nerve stimulator services. A new Category I code (42975) was also approved for Drug-Induced Sleep Endoscopy, which is the final procedure to determine which patients are appropriate for Inspire therapy. These new codes went into effect on January 1, 2022. With these approvals, a formal survey was conducted to determine the Medicare reimbursement levels assigned to each code and in November 2021 the final 2022



reimbursement rates were announced by the Centers for Medicare and Medicaid Services ("CMS"). The 2022 national average physician payments are approximately \$888 for implantation of a hypoglossal nerve stimulator and approximately \$115 for the DISE procedure. The 2022 rates of Medicare reimbursement to our hospital customers is approximately \$30,063, an increase of 2% over the 2021 rate. The ASC reimbursement rate for 2022 is approximately \$24,828, an increase of 2% over the 2021 rate.

We generated revenue of \$233.4 million, with a gross margin of 85.7% and a net loss of \$42.0 million, for the fiscal year ended December 31, 2021, compared to revenue of \$115.4 million, with a gross margin of 84.7% and a net loss of \$57.2 million, for the fiscal year ended December 31, 2020, and revenue of \$82.1 million, with a gross margin of 83.4% and a net loss of \$33.2 million, for the fiscal year ended December 31, 2019. Our accumulated deficit as of December 31, 2021 was \$279.4 million.

## Our Competitive Strengths

We believe the continued growth of our company will be driven by the following competitive strengths:

- ***First to market with an innovative, closed-loop, minimally invasive solution.*** We have developed the first and only FDA-approved neurostimulation technology that provides a safe and effective treatment for patients with moderate to severe OSA who have been confirmed to fail or cannot tolerate PAP treatments, such as CPAP. We received a PMA from the FDA in April 2014 for our Inspire therapy. Unlike CPAP, which is limited by low patient compliance primarily due to patient discomfort with the mask or device, our innovative, closed-loop, minimally invasive solution is designed to provide comfort and convenience, resulting in high compliance for patients with moderate to severe OSA. We believe we have a significant first mover advantage and momentum over future competitors, as physicians have treated more than 20,000 patients with Inspire therapy.
- ***Significant body of strong clinical data.*** We have developed a significant body of clinical data that demonstrates the safety and effectiveness, therapy adherence, and long-term sustained benefits of our Inspire therapy. The benefits of treatment with Inspire therapy have been consistent across seven sponsored and more than 70 independent clinical studies that evaluated several thousand patients, and have been highlighted in more than 190 peer-reviewed publications. Data reported in these clinical studies also demonstrated a high level of overall patient satisfaction. We believe this favorable data provides us with a significant competitive advantage and will continue to support increased adoption of our Inspire therapy.
- ***Holistic and targeted approach to market development and patient engagement.*** We have established a methodical approach to market development which centers on active engagement across three key stakeholders in the OSA treatment paradigm: physicians, sleep centers, and patients. Our sales force is focused on building long-lasting relationships with ENT physicians and sleep centers as we support physicians through all aspects of a case-from diagnosis to surgical support to patient follow-up. In addition, we are highlighting our compelling clinical data set and value proposition to increase awareness and adoption amongst referring physicians. We build upon this top-down approach with a strong direct-to-consumer marketing initiative that further drives demand through patient empowerment. This outreach helps to educate thousands of patients on our Inspire therapy. We are confident that this holistic approach to engagement across multiple constituents will continue to drive increased awareness of and demand for our Inspire therapy.
- ***Dedicated team focused on providing market access for patients and providers.*** We have a highly efficient approach to advance patients, once identified, to placement of the Inspire system. Our dedicated market access team helps patients and providers work with payors to secure prior authorization approvals in advance of initial treatment. In addition, this team proactively works with payors to establish positive coverage policies when needed by highlighting the compelling clinical data and the value of our Inspire therapy. This highly leverageable team has been successful in helping to secure reimbursement from hundreds of commercial payors to date, and positive coverage policies from most U.S. commercial payors, including virtually all large national payors.

- ***Strong research and development capabilities and comprehensive intellectual property portfolio.*** Our commitment to driving innovation has allowed us to achieve continuous, significant improvements of our Inspire therapy. For example, in February 2021, the FDA approved a new Inspire physician programmer system and in March 2021, the FDA approved an update to the system manual for the Inspire system to include an improved surgical implant procedure that eliminates one incision with a revised placement of the pressure sensing lead. In May 2021, we received certificates of conformity in Europe for the two-incision implant procedure which allowed us to affix the CE mark on our system. In December 2021, the FDA approved our new Bluetooth-enabled patient remote. We have a comprehensive patent portfolio to protect our intellectual property and technology, with rights as of December 31, 2021 to 44 issued U.S. patents, 34 issued foreign patents, 75 pending U.S. patent applications, and 63 pending foreign patent applications that cover aspects of our Inspire system and future product concepts.

## **Our Strategy**

Our goal is to be a global leader in providing clinically proven innovative solutions that improve sleep, quality of life, and health of patients with moderate to severe OSA. We believe the following strategies will play a critical role in achieving this goal and our future growth:

- ***Promote awareness among patients, ENT physicians, sleep centers, and referring physicians.*** We believe that many patients who have failed or cannot tolerate CPAP are unaware of our Inspire therapy as a safe and effective alternative treatment for moderate to severe OSA. We intend to continue to promote awareness of our therapy through training and educating ENT physicians, sleep centers, key opinion leaders, and various medical societies on the proven clinical benefits of Inspire therapy. In addition, we intend to continue to publish additional clinical data in various industry and scientific journals and online, and to present at various industry conferences. We also plan to continue building patient awareness through our direct-to-consumer marketing initiatives, which include paid online search, radio, social media, television, and online videos.
- ***Expand our sales and marketing organization to facilitate adoption of our Inspire therapy.*** We plan to expand our sales and marketing organization and seek to recruit and train exceptionally talented sales representatives in existing and new markets in the U.S. and in Europe to help facilitate further adoption and broaden awareness of our Inspire therapy. Our success to date in developing new markets has been primarily due to our ability to identify new regions with high volume medical centers, educate ENT and sleep physicians, help generate steady patient demand, and provide sufficient support staff to our sales representatives. We believe investing in a scalable, efficient direct sales force and continuing the development of our marketing efforts will help us broaden adoption of our Inspire therapy and drive revenue growth.
- ***Leverage our prior authorization model while we work with payors to broaden coverage.*** Our dedicated in-house market access team will continue to assist patients and physicians in obtaining prior authorization approvals from commercial payors for treatment with our Inspire therapy. In parallel, we are in active discussions with the commercial payors that have yet to establish positive coverage policies and modify existing policies when needed. We believe increased positive payor coverage policies and modifying existing positive coverage policies could expand patient access by reducing hurdles to treatment.
- ***Invest in research and development to drive innovation and expand indications.*** Our foundational commitment to driving innovation and improving patient lives fuels our desire for continuous product development. We intend to invest in existing and next generation technologies to further improve our products and clinical outcomes, optimize patient acceptance and comfort and broaden the patient population that can benefit from our Inspire therapy. Recent examples of our product innovation include the next generation of the Inspire neurostimulator, which is in development, and an active project to improve the physician programmer. We have launched a cloud-based patient management system called Inspire Cloud, which allows physicians to monitor patient compliance and more efficiently coordinate patient care, and in 2020, we launched the Inspire Sleep app for patients' smart phones. In December

2021, the FDA approved our new patient remote control which is Bluetooth® enabled. An example of our efforts to expand our label indications includes our clinical study that is evaluating the use of Inspire therapy in pediatric patients with Down syndrome. In April 2020, we received FDA approval for an expanded age-range for Inspire therapy to include 18 to 21 year old patients. In August 2020, the Australian Therapeutic Goods Administration approved Inspire therapy to treat moderate to severe OSA, and Japan's MLHW approved Inspire therapy to treat moderate to severe OSA in June 2018.

- **Further penetrate and expand into existing and new international markets.** We plan to establish and strengthen our presence internationally. Our goal is to further increase sales of our Inspire therapy in existing international markets in Europe, including Germany and the Netherlands, and in Japan. We plan to expand our reach to markets in new regions, such as Australia, Singapore, Hong Kong, South Korea and China. We plan to strategically invest in new markets based on our assessment of market size and opportunity and prospects for compelling reimbursement coding and coverage.

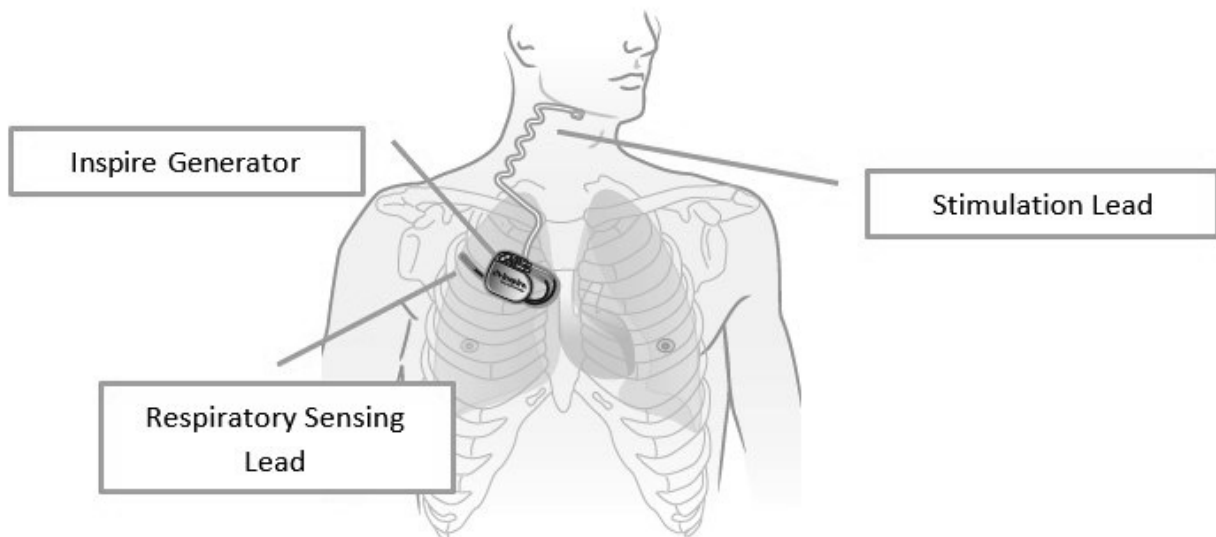
## Our Solution for OSA

### Overview of Inspire Therapy

Our proprietary Inspire system is the first and only FDA-approved closed-loop neurostimulation technology that provides a safe and effective treatment for moderate to severe OSA. Our Inspire system consists of a remote control and three implantable components:

- a pressure sensing lead, which detects when the patient is attempting to breathe;
- a neurostimulator, which houses the electronics and battery power for the device; and
- a stimulation lead, which delivers electrical stimulation to the hypoglossal nerve.

The image below depicts the location of the Inspire system under the patient's skin:



A pressure sensing lead is used to monitor the patient's breathing. Our proprietary algorithm tracks breathing patterns and the neurostimulator delivers electrical stimulation at the start of inspiration. This electrical stimulation of the hypoglossal nerve causes a slight forward movement of the back of the tongue that helps maintain an open airway, thereby preventing obstructive events and enabling the patient to inhale freely.

To receive the Inspire system, patients undergo a short outpatient surgical procedure, typically lasting about 90 minutes, during which the neurostimulator, sensing lead, and stimulation lead are implanted. The procedure is minimally invasive and performed with two small incisions. Patients typically recover quickly and are able to resume normal activities in just a few days. Initial activation of the system occurs 30 days after the implantation. After the initial activation, the patient is instructed to use the therapy each night by turning on their Inspire system before going to sleep using their remote control.

The following pictures depict the Inspire neurostimulator, shown with a quarter for scale, and the patient remote control, shown in hand for scale.



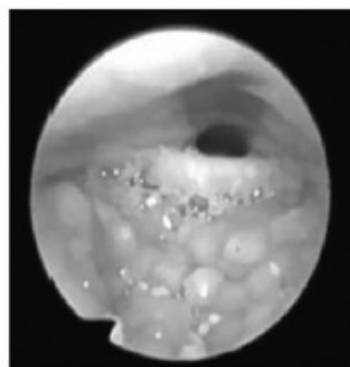
Patients turn their Inspire system on when they plan to go to sleep and turn it off when they awaken. The device has a programmed delay, typically 30 minutes, to allow patients to fall asleep naturally before the device activates. It then monitors the patient's breathing and delivers mild stimulation to the hypoglossal nerve at the start of the inspiratory phase, causing a slight forward movement at the back of the tongue to maintain an open airway during the inspiratory phase of respiration. The therapy is designed to provide stimulation for each breath to prevent obstructive events.

The following pictures depict the anatomy of a patient experiencing an OSA event. The patient's soft palate and the base of the patient's tongue are obstructing the patient's airway and limiting airflow to the lungs.

#### Obstructed Airway



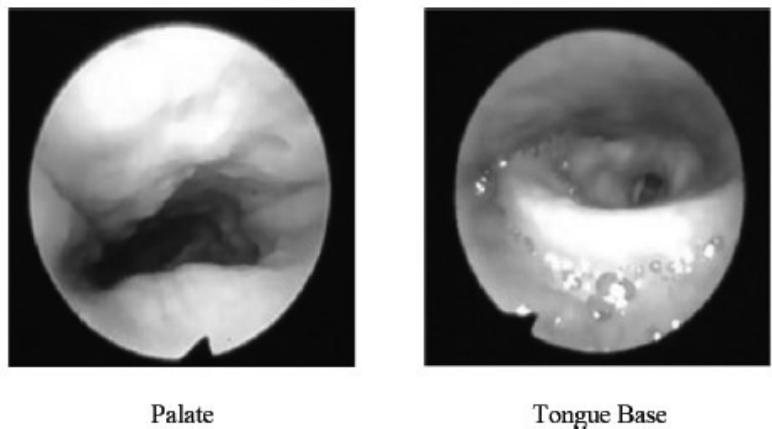
Palate



Tongue Base

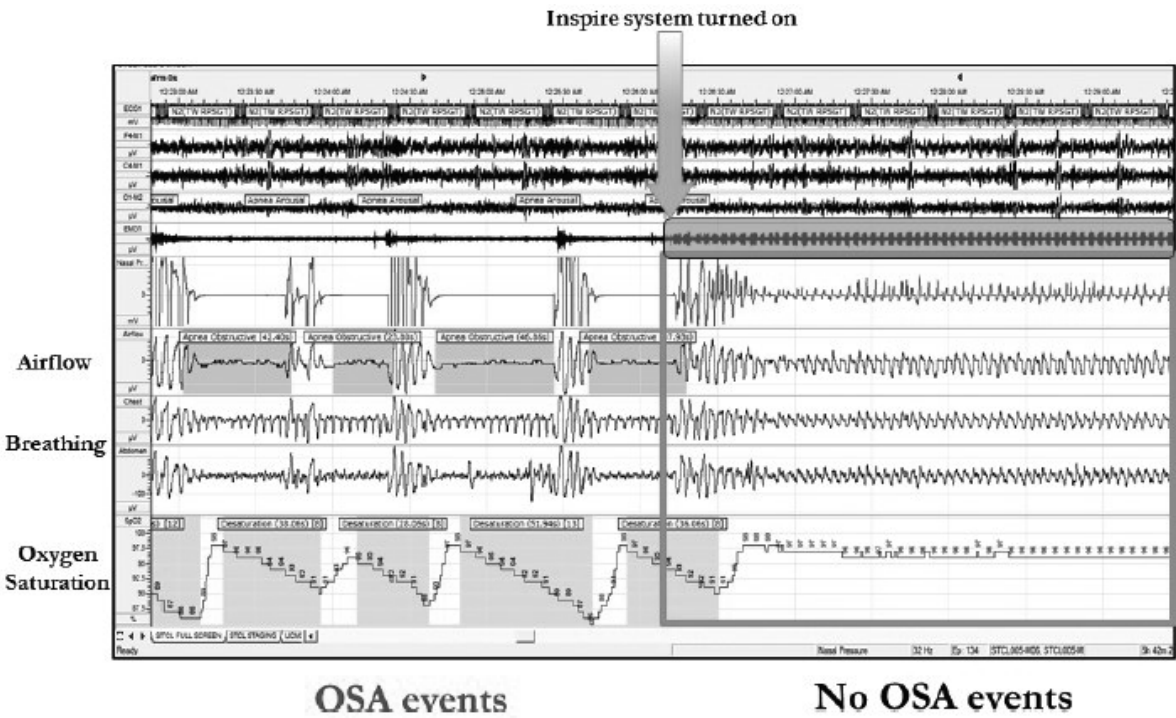
The following pictures depict the anatomy of the patient after mild stimulation of the hypoglossal nerve, which caused the patient's tongue to move forward slightly, opening the patient's airway and restoring airflow to the lungs.

Open Airway



The effectiveness of Inspire therapy to relieve OSA is objectively measured during a sleep study or polysomnogram. A sleep study records a patient's breathing, airflow, and blood oxygen levels before and after activating the device. Before activation, the patient experiences multiple periods of interrupted breathing, and oxygen levels repeatedly drop before the patient experiences a transient arousal that allows air intake. The polysomnogram below shows that after activating Inspire therapy, the patient exhibited a more regular breathing pattern, higher and more consistent blood oxygen levels, and fewer or no transient arousals.

Polysomnogram Before and After Activation of Inspire System



## ***Benefits of Inspire Therapy***

We believe our Inspire therapy overcomes many of the limitations of CPAP and other current treatments of moderate to severe OSA by providing the following key benefits:

- **Safe, effective, and durable treatment.** Results from our clinical trials provide compelling safety and efficacy data regarding the clinical benefits of Inspire therapy as many as five years after initial treatment. The results from our STAR trial, a five-year follow-up phase III pivotal trial, demonstrated an approximately 70% reduction in the median AHI from a baseline of 29.3 events per hour to 9.0 events per hour at 12 months following initial treatment. STAR trial follow-up has shown similar results to the initial data at 18 months, three years, and five years. At five years, median AHI remained low at 6.2 events per hour.
- **Closed-loop system.** The Inspire system uses a proprietary algorithm to continuously monitor a patient's breathing and provide electrical stimulation during the inspiratory phase, working with the body's natural actions to keep the airway open during the breathing cycle.
- **Comfortable and convenient therapy resulting in high patient satisfaction.** Data reported on the first 1,849 patients in our ongoing ADHERE patient registry, which we established to follow patients who have been implanted with an Inspire system, demonstrated that these patients used Inspire therapy an average of 5.6 hours per night an average of 12 months after initial treatment, with overall patient satisfaction reported to be at 91%.
- **Strong patient compliance.** Results from our STAR trial demonstrated that 80% of patients continue to use Inspire therapy on a nightly basis five years after initial treatment.
- **Minimally invasive outpatient procedure.** The Inspire system's implantable components are placed during an approximately 90-minute outpatient procedure. The procedure is minimally invasive and performed with two small incisions. Patients typically recover quickly and are able to resume normal activities within a few days.
- **Long-lasting solution.** Our Inspire system uses a battery designed to last approximately 11 years without charging or maintenance.

## ***Commercialization of Inspire Therapy***

In the U.S., before we can market a new medical device, or a new use of, new claim for or significant modification to an existing product, we must first receive FDA clearance or approval. We obtained PMA for our Inspire system in 2014. Additionally, we received a certificate of conformity for commercialization of our Inspire system in Europe in 2011 which allowed us to affix the CE mark on our device. Japan's MLHW approved Inspire therapy to treat moderate to severe OSA in June 2018 and was formally added to the Japan National Health Insurance Payment Listing in June 2021. In August 2020, the Australian Therapeutic Goods Administration approved Inspire therapy to treat moderate to severe OSA, and we are currently seeking reimbursement coverage in Australia.

To commercialize our Inspire system, both in the U.S. and Europe, we focus on physician and patient awareness and adoption of our Inspire therapy. To achieve this, our commercialization strategy primarily consists of our direct sales force engaging in sales efforts and promotional activities focused on ENT physicians and sleep centers and highlighting our compelling clinical data and value proposition. Our direct sales force utilizes strong direct-to-consumer marketing initiatives to create awareness of the benefits of our Inspire system. We intend to make significant investments building our sales and marketing organization by increasing the number of U.S. sales representatives and continuing our direct-to-consumer marketing efforts in existing and new markets throughout the U.S. and Europe.

In Japan, our commercialization approach is through our exclusive distribution partner, Japan Lifeline ("JLL"). JLL is responsible for local sales and promotional activities focused on ENT physicians, sleep centers, and community



awareness. We work closely with JLL to ensure a globally consistent approach and effective employee and customer training are in place.

In addition, a significant part of our commercialization effort consists of supporting our customers through the reimbursement process. Most commercial U.S. insurers now cover Inspire therapy. For those payors that do not have a positive policy, Inspire provides robust prior authorization support to our customers. Medicare also has positive coverage for all states, and we have a U.S. government contract for patients who are treated by the Veterans Health Administration.

## **Treatment with Inspire Therapy**

### ***Patient Selection***

Inspire therapy is indicated for patients with moderate to severe OSA (AHI of 15 to 65) who do not have significant central sleep apnea and do not have a complete concentric collapse of the airway at the soft palate level. Patients undergo a drug-induced sleep endoscopy performed by an ENT surgeon in order to confirm that they satisfy this anatomical requirement. In addition, patients in the U.S. and Japan must have been confirmed to fail or be unable to tolerate PAP treatments, such as CPAP, and be 18 years of age or older, though there are no similar requirements for patients in Europe. Patients who fail PAP are those that are not able to eliminate moderate to severe OSA despite PAP usage. Patients who cannot tolerate PAP treatments are those who either are unable to use PAP more than five nights per week for at least four hours per night, or who are unwilling to use PAP treatment.

### ***Implantation***

The Inspire system is implanted under general anesthesia through two small incisions. One incision is under the lower jaw, where the stimulation lead is attached around a distal branch of the hypoglossal nerve that is responsible for forward movement of the tongue. A second incision in the upper right chest below the clavicle is used to implant the neurostimulator, which houses all the electronics and battery power for the device, and a pressure sensing lead to monitor the breathing cycle. The functionality of the Inspire system is tested in the operating room to verify proper placement of the stimulation and pressure sensing leads. The wires for the electrodes are tunneled under the skin and the incisions are closed. The Inspire system is powered by an internal battery that is designed to last approximately 11 years without needing to be recharged. After this time, the neurostimulator is replaced during a simple outpatient procedure.

The implantation procedure is performed in an outpatient setting and surgery is completed in approximately 90 minutes. Patients may experience mild discomfort and swelling at the incision sites for a few days that is usually managed with over-the-counter pain medications. Patients can return home and resume a normal diet shortly after completion of the procedure and resume most daily activities within a few days. The only restriction on their activity is to avoid strenuous activities until the incisions have had time to heal.

### ***Activation***

Patients are allowed to heal for a month before the Inspire system is activated through a wireless connection to the device in the clinician's office. The initial activation is performed by the clinician using a programming tablet that is able to turn the system on as well as change various parameters such as the strength of the stimulating pulse, the sensitivity of the detection, the timing and length of the pulse, and which part of the stimulating electrode should be used. With the exception of pulse strength, the factory default settings are used in the majority of patients. The pulse strength is initially adjusted to the lowest level required to move the tongue out of the way without causing discomfort.

Patients receive a remote control that they use to turn their Inspire system on when they plan to go to sleep and to turn it off when they awaken. The device has a programmed delay, typically 30 minutes, to allow patients to fall asleep naturally before the device activates. It then delivers mild stimulation to the hypoglossal nerve, causing the tongue to move as the patient is inhaling. The remote enables patients to adjust the strength of the stimulation to optimize their therapy and comfort. The range of control given to patients is limited to avoid setting the strength of

the stimulation to an ineffective or excessively high level. Patients also have the ability to temporarily pause therapy if they awaken during the night.

## Clinical Results and Studies

A significant body of published clinical evidence, which includes seven sponsored and more than 70 independent clinical studies that evaluated several thousand patients, supports the safety and effectiveness of our Inspire therapy. The results of the STAR trial, our phase III pivotal clinical trial that served as the basis for the FDA approval of our PMA application, were published in the *New England Journal of Medicine*, and the results of additional clinical studies have been published in more than 190 peer-reviewed publications. We have established a global patient registry, which we refer to as our ADHERE patient registry, to collect data on safety, effectiveness, weekly usage, overall compliance, and satisfaction from patients who have been implanted with an Inspire system. The table below highlights key findings from certain of these studies and data from the first 1,849 patients in our ADHERE patient registry, including significant improvements in objective sleep measures and patient-reported quality of life measures, strong therapy compliance, and a favorable safety profile.

	STAR Trial <sup>(1)</sup>		German Post-Market Study <sup>(1)</sup>	ADHERE Patient Registry <sup>(1)</sup>	TJUH and UPMC Evaluation <sup>(2)</sup>
Number of Inspire therapy patients	124	97	56	1,849	97
Time following implantation	12 months	5 years	12 months	12 months	3 months
AHI—Baseline	29.3	29.3	28.6	33.0	35.6
AHI—Therapy	9.0	6.2	9.5	9.9	6.3
ESS—Baseline	11	11	13	11	11
ESS—Therapy	6	6	7	6	6
FOSQ—Baseline	14.6	14.6	13.7	*	*
FOSQ—Therapy	18.2	18.7	18.6	*	*
Therapy compliance	86% daily; 93% 5+ days weekly	80% daily	Average 39 hours per week; 89% ≥20 hours per week	Average 5.6 hours per night	Average >45 hours per week

\* Not measured

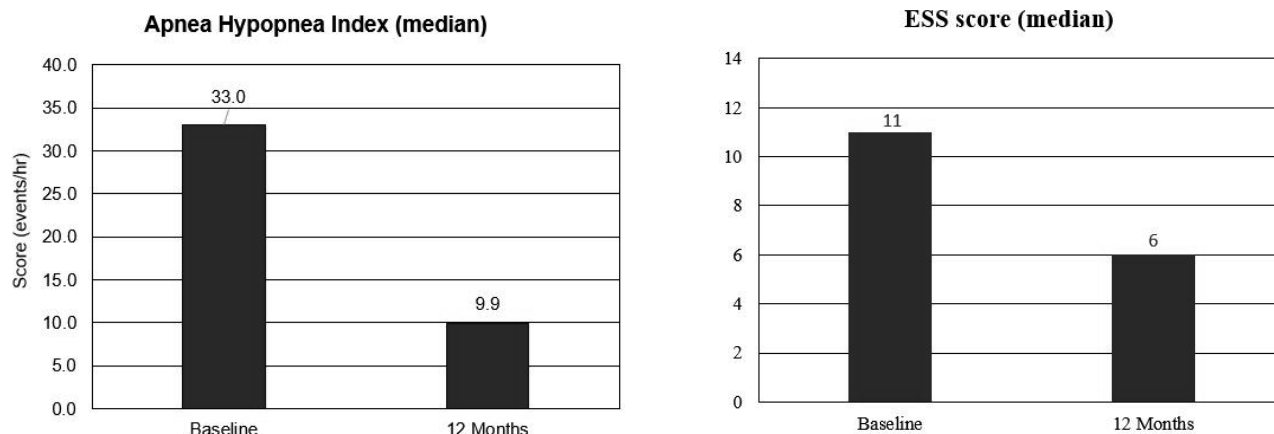
(1) Median results

(2) Thomas Jefferson University Hospital ("TJUH") and University of Pittsburgh Medical Center ("UPMC"). Mean results

## Select Recent Clinical Results and Studies

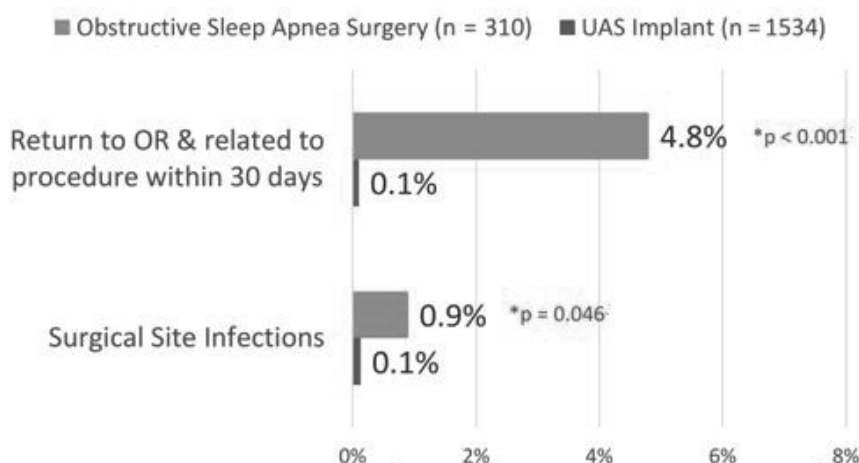
### ADHERE Patient Registry

We established our ADHERE patient registry to follow patients who have been implanted with an Inspire system, with a goal of collecting data on a group of at least 5,000 patients. Data gathered on the first 1,849 patients published in *Laryngoscope* in 2021 showed that patients used Inspire therapy an average of 5.6 hours per night when measured an average of 12 months after implantation. Median AHI was reduced from 33.0 events per hour to 9.9 events per hour and median ESS score improved from 11 to 6 over the same period. Overall satisfaction with Inspire therapy was reported by patients to be 91%, with 93% of patients reporting that they would choose the procedure again. In addition, 92% of patients reported a better experience than CPAP.



#### Comparison of Sleep Surgery Complication Rates versus Inspire Therapy

A 2021 paper published in *Otolaryngology – Open Journal* found that Inspire therapy had a lower complication rate and shorter hospital length of stay compared with sleep surgery, despite the fact that the Inspire population was older and had more co-morbidities. Combined with previous studies on sleep surgery outcomes, this demonstrates that Inspire therapy may be safer and more effective than sleep surgery in selected patients.



#### Upper Airway Surgery Improves Patient Symptoms Similarly as CPAP

In 2021, Cleveland Clinic published on the first comparison of patient-reported outcomes between Upper Airway Surgery ("UAS") and CPAP. These results showed that UAS and CPAP had similar degree of improvement of patient symptoms of sleepiness, daytime energy, depression, and insomnia. The results also showed that UAS patients were more likely to have a clinically meaningful improvement of symptoms than CPAP patients. This high degree of improvement was maintained through at least one-year in the UAS patients. In this first study comparing UAS versus CPAP patient symptoms, they conclude that this improvement is likely due to UAS's high ease of use, effectiveness, and high rates of adherence, recommending it as an alternative to CPAP.

#### Two-incision Procedure has Similar Outcomes as Three-incision Procedure

The new two-incision implant technique was compared to the legacy three-incision technique. This study of several hundred patients in a multi-center registry showed that the new two-incision technique is 39 minutes faster than the legacy technique (129 minutes vs. 86 minutes, p<0.001), while maintaining similarly low complication rates of 1%, and equivalent outcomes of reductions in AHI and Epworth Sleepiness Scale. Thus, the new two-

incision technique is faster, without compromising safety and efficacy, saving operative time for physicians and enabling shorter recovery time for patients.

## **Sales and Marketing**

We have established a methodical approach to market development which centers on active engagement across three key stakeholders in the OSA treatment paradigm: patients, physicians, and sleep centers.

We sell our Inspire system through a direct sales force that primarily targets ENT physicians and sleep centers in the U.S. and Europe, and through a distributor in Japan. The implant procedure for our Inspire therapy is typically performed by an ENT physician or in some cases by neurosurgeons. We also focus on sleep centers because they diagnose and manage large volumes of patients with sleep apnea and are often an important referral base for ENT physicians. In addition, because OSA is sometimes diagnosed during other procedures, we have developed programs to help educate general practitioners and specialists in other fields, such as cardiovascular surgeons, electrophysiologists, and dentists, regarding our Inspire therapy.

We have 157 sales territories in the U.S. and 12 in Europe. We seek to recruit sales representatives with strong sales backgrounds, direct experience developing markets with new technologies, and core knowledge of medical device coding, reimbursement, and the prior authorization process. In Japan, we rely on our distribution partner for local sales and promotional activities.

We also utilize direct communication channels to inform and educate patients about Inspire therapy and to enable them to connect with active clinical sites that offer our Inspire systems. Our primary methods of patient outreach are Facebook, Google ad placements, radio advertisements (either local or satellite), and television advertisements. The objective of this outreach is to bring patients to our website, where they can find educational materials and videos on sleep apnea and the use and benefits of our Inspire therapy, contact information for physicians and clinical sites, and information regarding community awareness events.

We believe our patient outreach efforts have been effective in bringing potential patients to our website and facilitating contact with our clinical sites. During 2021, we had over 7.3 million visits to our website, of which 410,000 used our "Find a Doctor" function to find a physician in their area. This generated more than 95,000 contacts with physicians throughout the year and approximately 33,000 visitors to our website signed up to attend a local or online Inspire informational event.

In 2020, we launched the Inspire Sleep app for patients' smart phones. This app is an educational tool for patients and also interfaces with the Inspire Cloud to allow physicians to collect clinical data from patients directly. We continue to enhance the functionality of this app as part of our overall digital platform development. During 2020 and 2021, over 26,000 copies of the app were downloaded to smartphones.

## ***Commercial Activities Outside of the U.S.***

Our general practice is to limit commercial investments in countries until such time as there is a determined reimbursement pathway. We have 12 sales territories in Europe, 8 of which are located in Germany, and we sell our products through a distributor in Japan. We provide consistent training in Europe and Japan as is conducted in the U.S. and have established a support team in Europe for patient outreach and education, implant support, and device programming. In Japan we assist our distribution partner with patient outreach and education initiatives. We expect to continue to scale our commercial activities in Europe as we continue to develop country-wide reimbursement in additional markets. We continue to work on the reimbursement process in Australia and hope to commercialize Inspire therapy there.

## **Third-Party Reimbursement**

Our market access team is responsible for all of our reimbursement processes and initiatives. Our team includes 20 professionals who are focused on all key aspects of reimbursement, which include coding, payment, coverage, and prior authorization.

## ***Coding and Payment***

In the U.S., we sell our products to hospitals and ASCs. These customers in turn bill various third-party payors, such as commercial payors and Medicare, for the cost required to treat each patient.

Third-party payors require physicians and hospitals to identify the service for which they are seeking reimbursement by using CPT codes, which are created and maintained by the AMA. The procedure performed to implant our device was previously described for billing purposes using a Category I CPT code (64568), which was used in conjunction with a temporary Category III CPT code (+0466T). At the October 2020 AMA CPT Editorial Panel meeting, the AMA approved the creation of new Category I CPT codes (64582, 64583, and 64584) to separately identify hypoglossal nerve stimulator services. A new Category I code (42975) was also approved for DISE, which is the final procedure to determine which patients are appropriate for Inspire therapy. These new codes went into effect on January 1, 2022.

Physician reimbursement under Medicare is based on a defined fee schedule, the Physician Fee Schedule, through which payment amounts are determined by the relative values of the professional service rendered. Medicare provides reimbursement to our hospital customers under the hospital outpatient prospective payment system ("HOPPS") which provides bundled amounts generally intended to reimburse the hospital for all facility costs related to procedures performed in the hospital outpatient setting. Under the HOPPS, the 2022 national average Medicare payment to the hospital for this procedure is \$30,063, which is intended to cover the hospitals' costs for the device and the implantation procedure. The 2022 ASC reimbursement rate is \$24,828. The physician's professional services are reimbursed separately under the Medicare Physician Fee Schedule. The 2022 national average physician fee for the implant (CPT code 64582) is \$888. The 2022 average reimbursement rate for the DISE procedures is \$115.

Reimbursement rates from commercial payors vary depending on the procedure performed, the commercial payor, contract terms, and other factors.

## ***Commercial Payor and Government Program Coverage***

A core pillar of our reimbursement strategy involves broadening our third-party payor coverage when possible. We continue to have active discussions with commercial payors to establish new and modify existing positive coverage policies by highlighting our compelling and robust clinical data, increased patient demand, and support from leading medical societies and key opinion leaders. We have been successful in obtaining prior authorization approvals from most commercial payors for the Inspire device and procedure. Historically, commercial payors approve prior authorization requests for approximately 70% of Inspire implants in the U.S. We estimate that the majority of patients who meet the FDA indication for Inspire therapy are covered by commercial insurance companies and we have secured coverage policies with virtually all major national commercial payors.

A number of small regional commercial payors currently consider hypoglossal nerve stimulation to be investigational, including procedures involving the Inspire system, and therefore do not have formal coverage policies, which presents an opportunity for incremental positive coverage development.

All seven MACs published final policies in 2020 that provide coverage of Inspire therapy when certain coverage criteria are met. Medicare beneficiaries have historically accounted for approximately 25% of all Inspire system implantations in the U.S. In addition, we have a contract with the U.S. government that covers implantations of our Inspire system performed in Veterans Affairs and military hospitals, which account for approximately 5% of all Inspire system implantations historically in the U.S.

## ***Prior Authorization Approval Process***

A second pillar of our reimbursement strategy includes leveraging our market access team to assist patients and physicians in obtaining appropriate prior authorization approvals in advance of treatment on a case-by-case basis where positive coverage policies currently do not exist. We believe our market access team is highly effective in working with patients and physicians to obtain prior authorizations for our Inspire system including assisting with

the appeals process. We have received multiple prior authorization approvals from all of the largest commercial payors, for example Anthem, Cigna, Blue Cross Blue Shield, United Healthcare, and Humana.

We believe we will continue to benefit from this efficient prior authorization process in the near-term and in the longer-term by expanding positive coverage policies. We intend to expand our market access team and increase the number of annual patient submissions as we grow our operations.

### ***Reimbursement Outside of the U.S.***

In Germany, the Institut für das Entgeltsystem im Krankenhaus ("InEK"), the German federal reimbursement agency, fully integrated Inspire therapy into the German hospital reimbursement system ("G-DRG"), effective January 1, 2021. The integration was done under conditions of a Zusatzentgelt (a supplemental fee or add-on payment scheme). The reimbursement for the procedure has two parts, a G-DRG fee plus an add-on payment called a ZE fee, in this case, ZE2021-187 Neurostimulatoren zur Hypoglossusnerv-Stimulation (Neurostimulators for hypoglossal nerve stimulation). The amount for the G-DRG is calculated annually by the InEK based on empirical cost data supplied by reference hospitals. The ZE fee is negotiated locally with each hospital, paid additionally and usually covers device fees for the Inspire system, additional procedural equipment, and service costs that are not included within the G-DRG. Before January 1, 2021, Germany's reimbursement for the Inspire procedure was provided through Neue Untersuchungs-und Behandlungsmethoden ("NUB") Status 1 coverage. The NUB process allows for the introduction of new and innovative medical devices prior to reaching reimbursement eligibility and provides for a supplemental payment for new technologies in the German reimbursement system. NUB Status 1 is the highest of four levels and allowed for full reimbursement for our Inspire system. NUM reimbursement requires the negotiation of a respective agreement with the treating hospital and applies for a limited term. Under NUB Status 1, payors at these hospitals were obligated to cover the gaps in treatment costs for the Inspire system.

In the Netherlands, the Zorginstituut, or the Dutch Health Institute, has approved Inspire therapy for conditional reimbursement within the basic package of all payors. This allows patients to receive the Inspire system in certain hospitals and within approved indications set by the Zorginstituut.

In Switzerland, Inspire therapy is covered by the public health insurance and hospitals receive reimbursement from the Swiss Diagnosis Related Groups ("DRG") system. This allows patients to receive the Inspire system in certain hospitals and within the indication covered by the Swiss DRG.

In Japan, Inspire therapy is paid for by Japan's National Health Insurance system, following coverage requirements established by the Ministry of Health, Labour and Welfare.

Reimbursement in other countries can often be established through a combination of private (commercial insurance) and public funding sources, or at the hospital level through innovation budgets. We continue to pursue permanent reimbursement in target markets across the Asia Pacific, including Australia, and Europe, including Belgium, France, Spain, the United Kingdom, and the Nordic region.

## **Research and Development**

### ***Product Evolution and Next Generation Products***

The first Inspire device was developed by Medtronic Inc. (now Medtronic Public Limited Company), or Medtronic, in the early 1990s as a radio frequency controlled device that required an external apparatus to deliver electrical stimulation to the hypoglossal nerve. The first fully implantable, respiration-sensing, closed-loop Inspire system was developed shortly thereafter. Based on the initial clinical trial results, which were published in 2001, Medtronic began developing what became known as our Inspire II system, introducing a new, more durable stimulation lead and lower-power neurostimulator, and relocating the respiratory sensing lead to between the intercostal muscle layers.

After our November 2007 inception and contemporaneous spin-off of the Inspire business from Medtronic, our primary focus was to requalify the Inspire II system and resume clinical trial activity. We completed a phase I



feasibility trial along with a phase II dosing or patient selection trial in 2009. In 2011, we began our phase III pivotal STAR trial. The STAR trial was completed and published in the *New England Journal of Medicine* in January 2014 and we received PMA in April 2014. Additionally, we received a CE mark for commercialization in Europe in 2011.

We continue to invest in advancing our Inspire system with the goal of providing patients more effective and less invasive therapy for OSA. In 2017, we released the Inspire IV neurostimulator, which is 40% smaller than the previous version while maintaining approximately 11 years of battery life. Patients with this version of the Inspire system are now able to undergo an MRI scan of the head or extremities. The Inspire IV device was launched in the U.S. in July 2017, and in Europe in the second quarter of 2018.

Our newly designed respiratory sensing lead was developed in 2017 and was approved by the FDA in January 2019. This lead was introduced in Europe in October 2018 and in the U.S. in March 2019. The lead has a significantly smaller profile, including a reduction in diameter of 49 percent as compared to the prior design. This smaller profile is intended to simplify surgical placement along with several other features to enable easier implanting, and potentially reduce surgical time for an ENT surgeon.

Our next generation of the Inspire neurostimulator is in development, and we also have an active project to improve the physician programmer. Our Bluetooth® enabled patient remote control was approved by the FDA in December 2021. We have launched a cloud-based patient management system called Inspire Cloud, which allows physicians to monitor patient compliance and more efficiently coordinate patient care. In 2020, we launched the Inspire Sleep app for patients' smart phones. The first version of the app was an educational tool, and the second version interfaces with the Inspire Cloud and allows physicians to collect clinical data from patients directly. We continue to enhance the functionality of this app as part of our overall digital platform development.

The Inspire Cloud and our app are initial steps in establishing interconnectivity between the patient and their healthcare provider with a long-term plan to improve outcomes by tracking patient activity and adherence, and monitoring for any issues with device use.

### ***Additional Indications***

We have sought and continue to seek to expand the approved indications for our Inspire therapy. For instance, in January 2017, the FDA approved a PMA supplement expanding the indicated AHI range for our Inspire therapy from 20 to 65 events per hour to 15 to 65 events per hour. In April 2020, we received FDA approval for an expanded age-range of patients who can be treated with Inspire therapy, which now includes 18 to 21-year old patients.

Our research and development team focuses on the products currently under development, including our clinical trials, as well as feasibility studies in which we are evaluating different design configurations to enhance product functionality for future generations of the Inspire system.

### **Competition**

Our industry is subject to rapid change from the introduction of new products and technologies and other activities of industry participants. We compete as a second-line therapy in the OSA treatment market for patients with moderate to severe OSA.

We consider our primary competition to be other neurostimulation technologies designed to treat OSA, though we are currently the only such technology approved for commercialization in the U.S. by the FDA. Outside the U.S., we compete with LivaNova and Nyxoah. LivaNova, which markets an open-loop neurostimulation device, is currently conducting clinical trials of its device in the U.S. Nyxoah markets an open-loop bilateral hypoglossal nerve stimulation device in certain countries outside the U.S. and is conducting its first pivotal trial as it seeks FDA approval in the U.S. We believe other emerging businesses are in the early stages of developing neurostimulation devices or early-stage pharmaceutical approaches.

We also compete, both within and outside of the U.S., with invasive surgical treatment options such as UPPP, maxillomandibular advancement ("MMA") and robotic tongue reduction surgery, and, to a lesser extent, oral appliances, which are primarily used in the treatment of mild to moderate OSA. We do not believe we directly compete with CPAP or other types of PAP devices because in the U.S., Inspire therapy is only indicated for patients who have been confirmed to fail or cannot tolerate PAP treatments, such as CPAP.

We believe that the primary competitive factors in the OSA treatment market are:

- company, product, and brand recognition;
- product safety, efficacy, reliability, and durability;
- quality and volume of clinical data;
- effective marketing to and education of patients, physicians, and sleep centers;
- product ease of use and patient comfort;
- sales force experience and access;
- product support and service;
- technological innovation, product enhancements, and speed of innovation;
- pricing and revenue strategies;
- procedure costs to patients;
- effectiveness of reimbursement teams and strategies; and
- dedicated practice development and clinical training teams.

Some other OSA treatments against which we compete, such as oral appliances, MMA, and UPPP, have a greater penetration into the OSA treatment market. Oral appliances and some other surgical treatments are better known to ENT physicians, sleep centers and the other physicians on whom we rely for referrals, but we believe physician awareness of our Inspire therapy is increasing.

We also compete with other medical technology companies to recruit and retain qualified sales, training and other personnel, including members of our in-house prior authorization team.

### **Intellectual Property**

We rely on a combination of patent, copyright, trademark and trade secret laws, and confidentiality and invention assignment agreements to protect our intellectual property rights. As of December 31, 2021, we had rights to 44 issued U.S. patents, which will expire between 2022 and 2037 assuming all required fees are paid, 75 pending U.S. patent applications, 34 issued foreign patents, and 63 pending foreign patent applications. Our patents cover aspects of our current Inspire system and future product concepts. Some of the issued foreign patents and pending foreign patent applications preserve an opportunity to pursue patent rights in multiple countries.

There is no active patent litigation involving any of our patents and we have not received any notices of patent infringement.

As of December 31, 2021, we had 122 pending and registered trademark filings worldwide, some of which may apply to multiple countries.

We also rely, in part, upon unpatented trade secrets, know-how and continuing technological innovation, and may in the future rely upon licensing opportunities, to develop and maintain our competitive position. We protect our proprietary rights through a variety of methods, including confidentiality and assignment agreements with suppliers, employees, consultants, and others who may have access to our proprietary information.

Our pending patent applications may not result in issued patents, and we cannot ensure that any current or subsequently issued patents will adequately protect our products or provide us with any competitive advantage. While there is no active litigation involving any of our patents or other intellectual property rights and we have not received any notices of patent infringement, we may be required to enforce or defend our intellectual property rights against third parties in the future. See Part I., "Item 1A. Risk Factors — Risks Related to Intellectual Property Matters" for additional information regarding these and other risks related to our intellectual property portfolio and their potential effect on us.

### **License Agreement with Medtronic**

In November 2007, we entered into an assignment and license agreement with Medtronic ("the Assignment and License Agreement") pursuant to which Medtronic assigned certain patents and trademarks to us and granted to us a worldwide, royalty-free license to certain other patents and technical information to make, use, import and sell products, and to practice methods in the field of electrical stimulation of the upper airway for the treatment of OSA ("the Field"). We share co-exclusive rights with Medtronic under this license; however, Medtronic may not exercise its rights unless we make an assignment for the benefit of our creditors, file or have filed against us a bankruptcy petition or go into receivership. We also granted to Medtronic certain worldwide, royalty-free, exclusive licenses to the patents Medtronic assigned to us, as well as other intellectual property (including but not limited to Technical Information (as defined in the Assignment and License Agreement)) that applies to a device and methods with certain specifications for use in the Field, to make, use, import and sell products, and to practice methods outside of the Field. The licenses granted are perpetual and irrevocable.

### **Manufacturing and Supply**

We rely on third-party suppliers to manufacture our Inspire system and its components. Outsourcing manufacturing reduces our need for capital investment and reduces operational expense. Additionally, outsourcing provides expertise and capacity necessary to scale up or down based on demand for our Inspire system. We select our suppliers to ensure that our Inspire system and its components are safe and effective, adhere to all applicable regulations, are of the highest quality, and meet our supply needs. We employ a rigorous supplier assessment, qualification, and selection process targeted to suppliers that meet the requirements of the FDA and the International Organization for Standardization and quality standards supported by internal policies and procedures. Our quality assurance process monitors and maintains supplier performance through qualification and periodic supplier reviews and audits.

Certain components used in our Inspire system are supplied by single-source suppliers. Our suppliers manufacture the components they produce for us and test our components and devices to our specifications. We intend to maintain sufficient levels of inventory to enable us to continue our operations while we obtain another supplier in the event that one or more of our single-source suppliers were to encounter a delay in supply or end supply.

We have experienced and continue to experience some minor supply disruptions during the COVID pandemic, but have managed to avoid any significant supply and inventory issues.

### **Government Regulation**

Our products and our operations are subject to extensive regulation by the FDA and other federal and state authorities in the U.S., as well as comparable authorities in the European Economic Area ("EEA") and Japan, and in Australia (where our products are approved for sale but where we have not yet commercialized them). In the U.S., our products are subject to regulation as medical devices under the Federal Food, Drug, and Cosmetic Act ("FDCA") as implemented and enforced by the FDA. The FDA regulates the development, design, non-clinical and clinical research, manufacturing, safety, efficacy, labeling, packaging, storage, installation, servicing,

recordkeeping, premarket clearance or approval, import, export, adverse event reporting, advertising, promotion, marketing and distribution, and import and export of medical devices to ensure that medical devices distributed domestically are safe and effective for their intended uses and otherwise meet the requirements of the FDCA.

In addition to U.S. regulations, we are subject to a variety of regulations in the EEA governing clinical trials and the commercial sales and distribution of our products. Whether or not we have or are required to obtain FDA clearance or approval for a product, we will be required to obtain authorization before commencing clinical trials and to obtain marketing authorization or approval of our products under the comparable regulatory authorities of countries outside of the U.S. before we can commence clinical trials or commercialize our products in those countries. The approval process varies from country to country and the time may be longer or shorter than that required for FDA clearance or approval.

### ***FDA Premarket Clearance and Approval Requirements***

Unless an exemption applies, each medical device commercially distributed in the U.S. requires either FDA clearance of a 510(k) premarket notification or PMA. Under the FDCA, medical devices are classified into one of three classes—Class I, Class II, or Class III—depending on the degree of risk associated with each medical device and the extent of manufacturer and regulatory control needed to ensure its safety and effectiveness. Class I includes devices with the lowest risk to the patient and are those for which safety and effectiveness can be assured by adherence to the FDA's General Controls for medical devices, which include compliance with the applicable portions of the FDA's Quality System Regulation ("QSR"), facility registration and product listing, reporting of adverse medical events, and truthful and non-misleading labeling, advertising, and promotional materials. Class II devices are subject to the FDA's General Controls, and special controls as deemed necessary by the FDA to ensure the safety and effectiveness of the device. While most Class I devices are exempt from the 510(k) premarket notification requirement, manufacturers of most Class II devices are required to submit to the FDA a premarket notification under Section 510(k) of the FDCA requesting permission to commercially distribute the device. The FDA's permission to commercially distribute a device subject to a 510(k) premarket notification is generally known as 510(k) clearance. Under the 510(k) process, the manufacturer must submit to the FDA a premarket notification demonstrating that the device is "substantially equivalent" to either a device that was legally marketed prior to May 28, 1976, the date upon which the Medical Device Amendments of 1976 were enacted, or another legally marketed device that was cleared through the 510(k) process.

Devices deemed by the FDA to pose the greatest risks, such as life-sustaining, life-supporting or some implantable devices, or devices that have a new intended use, or use advanced technology that is not substantially equivalent to that of a legally marketed device, are placed in Class III, requiring approval of a PMA. Some pre-amendment devices are unclassified, but are subject to the FDA's premarket notification and clearance process in order to be commercially distributed.

Our currently marketed Inspire products are Class III devices which have received PMA.

### ***PMA Pathway***

In a PMA, the manufacturer must demonstrate that the device is safe and effective, and the PMA must be supported by extensive data, including data from preclinical studies and human clinical trials. The PMA must also contain a full description of the device and its components, a full description of the methods, facilities and controls used for manufacturing, and proposed labeling. Following receipt of a PMA, the FDA determines whether the application is sufficiently complete to permit a substantive review. If the FDA accepts the application for review, it has 180 days under the FDCA to complete its review of a PMA, although in practice, the FDA's review often takes significantly longer, and can take up to several years. An advisory panel of experts from outside the FDA may be convened to review and evaluate the application and provide recommendations to the FDA as to the approvability of the device. The FDA may or may not accept the panel's recommendation. In addition, the FDA will generally conduct a preapproval inspection of the applicant or its third-party manufacturers' or suppliers' manufacturing facility or facilities to ensure compliance with the QSR.

The FDA will approve the new device for commercial distribution if it determines that the data and information in the PMA constitute valid scientific evidence and that there is reasonable assurance that the device is safe and

effective for its intended use(s). The FDA may approve a PMA with post-approval conditions intended to ensure the safety and effectiveness of the device, including, among other things, restrictions on labeling, promotion, sale and distribution, and collection of long-term follow-up data from patients in the clinical study that supported PMA or requirements to conduct additional clinical studies post-approval. The FDA may condition PMA on some form of post-market surveillance when deemed necessary to protect the public health or to provide additional safety and efficacy data for the device in a larger population or for a longer period of use. In such cases, the manufacturer might be required to follow certain patient groups for a number of years and to make periodic reports to the FDA on the clinical status of those patients. Failure to comply with the conditions of approval can result in material adverse enforcement action, including withdrawal of the approval.

Certain changes to an approved device, such as changes in manufacturing facilities, methods, or quality control procedures, or changes in the design performance specifications, which affect the safety or effectiveness of the device, require submission of a PMA supplement. PMA supplements often require submission of the same type of information as a PMA, except that the supplement is limited to information needed to support any changes from the device covered by the original PMA and may not require as extensive clinical data or the convening of an advisory panel. Certain other changes to an approved device require the submission of a new PMA, such as when the design change causes a different intended use, mode of operation, and technical basis of operation, or when the design change is so significant that a new generation of the device will be developed, and the data that were submitted with the original PMA are not applicable for the change in demonstrating a reasonable assurance of safety and effectiveness.

### ***Clinical Trials***

Clinical trials are almost always required to support a PMA and are sometimes required to support a 510(k) submission. All clinical investigations of investigational devices to determine safety and effectiveness must be conducted in accordance with the FDA's investigational device exemption ("IDE") regulations which govern investigational device labeling, prohibit promotion of the investigational device, and specify an array of recordkeeping, reporting and monitoring responsibilities of study sponsors and study investigators. If the device presents a "significant risk" to human health, as defined by the FDA, the FDA requires the device sponsor to submit an IDE application to the FDA, which must become effective prior to commencing human clinical trials. A significant risk device is one that presents a potential for serious risk to the health, safety or welfare of a patient and either is implanted, used in supporting or sustaining human life, substantially important in diagnosing, curing, mitigating or treating disease or otherwise preventing impairment of human health, or otherwise presents a potential for serious risk to a subject. An IDE application must be supported by appropriate data, such as animal and laboratory test results, showing that it is safe to test the device in humans and that the testing protocol is scientifically sound. The IDE will automatically become effective 30 days after receipt by the FDA unless the FDA notifies the company that the investigation may not begin. If the FDA determines that there are deficiencies or other concerns with an IDE for which it requires modification, the FDA may permit a clinical trial to proceed under a conditional approval. An IDE supplement must be submitted to, and approved by, the FDA before a sponsor or investigator may make a change to the investigational plan that may affect its scientific soundness, study plan or the rights, safety or welfare of human subjects.

In addition, the study must be approved by, and conducted under the oversight of, an Institutional Review Board ("IRB") for each clinical site. The IRB is responsible for the initial and continuing review of the IDE, and may pose additional requirements for the conduct of the study. If an IDE application is approved by the FDA and one or more IRBs, human clinical trials may begin at a specific number of investigational sites with a specific number of patients, as approved by the FDA. If the device presents a non-significant risk to the patient, a sponsor may begin the clinical trial after obtaining approval for the trial by one or more IRBs without separate approval from the FDA, but must still follow abbreviated IDE requirements, such as monitoring the investigation, ensuring that the investigators obtain informed consent, and labeling and record-keeping requirements.

During a study, the sponsor is required to comply with the applicable FDA requirements, including, for example, trial monitoring, selecting clinical investigators and providing them with the investigational plan, ensuring IRB review, adverse event reporting, record keeping, and prohibitions on the promotion of investigational devices or on making safety or effectiveness claims for them. The clinical investigators in the clinical study are also subject to FDA regulations and must obtain patient informed consent, rigorously follow the investigational plan and study

protocol, control the disposition of the investigational device, and comply with all reporting and recordkeeping requirements. Additionally, after a trial begins, we, the FDA or the IRB could suspend or terminate a clinical trial at any time for various reasons, including a belief that the risks to study subjects outweigh the anticipated benefits.

### ***Post-market Regulation***

After a device is cleared or approved for marketing, numerous and pervasive regulatory requirements continue to apply. These include:

- establishment registration and device listing with the FDA;
- QSR requirements, which require manufacturers, including third-party manufacturers, to follow stringent design, testing, control, documentation, and other quality assurance procedures during all aspects of the design and manufacturing process;
- labeling and marketing regulations, which require that promotion is truthful, not misleading, fairly balanced, provide adequate directions for use, and that all claims are substantiated, and also prohibit the promotion of products for unapproved or “off-label” uses and impose other restrictions on labeling; FDA guidance on off-label dissemination of information and responding to unsolicited requests for information;
- clearance or approval of product modifications to 510(k)-cleared devices that could significantly affect safety or effectiveness or that would constitute a major change in intended use of one of our cleared devices, or approval of a supplement for certain modifications to PMA devices;
- medical device reporting regulations, which require that a manufacturer report to the FDA if a device it markets may have caused or contributed to a death or serious injury, or has malfunctioned and the device or a similar device that it markets would be likely to cause or contribute to a death or serious injury, if the malfunction were to recur;
- correction, removal, and recall reporting regulations, which require that manufacturers report to the FDA field corrections and product recalls or removals if undertaken to reduce a risk to health posed by the device or to remedy a violation of the FDCA that may present a risk to health;
- complying with the new federal law and regulations requiring Unique Device Identifiers (UDI) on devices and also requiring the submission of certain information about each device to the FDA’s Global Unique Device Identification Database (GUDID);
- the FDA’s recall authority, whereby the agency can order device manufacturers to recall from the market a product that is in violation of governing laws and regulations; and
- post-market surveillance activities and regulations, which apply when deemed by the FDA to be necessary to protect the public health or to provide additional safety and effectiveness data for the device.

Our manufacturing processes are required to comply with the applicable portions of the QSR, which cover the methods and the facilities, controls for the design, manufacture, testing, production, processes, controls, quality assurance, labeling, packaging, distribution, installation, and servicing of finished devices intended for human use. The QSR also requires, among other things, maintenance of a device master file, device history file, and complaint files. As a manufacturer, our facilities, records, and manufacturing processes are subject to periodic scheduled or unscheduled inspections by the FDA. Our failure to maintain compliance with the QSR or other applicable regulatory requirements could result in the shut-down of, or restrictions on, our manufacturing operations and the recall or seizure of our products. The discovery of previously unknown problems with any of our products, including unanticipated adverse events or adverse events of increasing severity or frequency, whether resulting from the use of the device within the scope of its clearance or off-label by a physician in the practice of medicine, could result in restrictions on the device, including the removal of the product from the market or voluntary or mandatory device recalls.



The FDA has broad regulatory compliance and enforcement powers. If the FDA determines that we failed to comply with applicable regulatory requirements, it can take a variety of compliance or enforcement actions, which may result in any of the following sanctions:

- warning letters, untitled letters, fines, injunctions, consent decrees, and civil penalties;
- recalls, withdrawals, or administrative detention or seizure of our products;
- operating restrictions or partial suspension or total shutdown of production;
- refusing or delaying requests for 510(k) marketing clearance or PMAs of new products or modified products;
- withdrawing 510(k) clearances or PMAs that have already been granted;
- refusal to grant export or import approvals for our products; or
- criminal prosecution.

### ***Foreign Regulation***

In order for us to market our products in countries outside the U.S., we must obtain regulatory approvals or certifications and comply with extensive product and quality system regulations in other countries. These regulations, including the requirements for approvals, clearance or certifications and the time required for regulatory review, vary from country to country. Some countries have regulatory review processes that are substantially longer than U.S. processes. Failure to obtain regulatory approval or certification in a timely manner and meet all of the local requirements including language and specific safety standards in any foreign country in which we plan to market our products could prevent us from marketing products in such countries or subject us to sanctions and fines.

### ***Regulation of Medical Devices in the European Union***

The European Union ("EU") has adopted specific directives and regulations regulating the design, manufacture, clinical investigation, conformity assessment, labeling and adverse event reporting for medical devices.

Until May 25, 2021, medical devices were regulated by Council Directive 93/42/EEC (the "EU Medical Devices Directive"), and Directive 90/385/EEC ("AIMDD") which have been repealed and replaced by Regulation (EU) No 2017/745 (the "EU Medical Devices Regulation"). Our current certificates have been granted under the EU Medical Devices Directive and the AIMDD whose regime is described below. However, as of May 26, 2021, some of the EU Medical Devices Regulation requirements apply in place of the corresponding requirements of the EU Medical Devices Directive and the AIMDD with regard to registration of economic operators and of devices, post-market surveillance and vigilance requirements. Pursuing marketing of medical devices in the EU will notably require that our devices be certified under the new regime set forth in the EU Medical Devices Regulation when our current certificates expire.

#### ***Medical Devices Directive***

In the EU, there is currently no premarket government review of medical devices. However, all medical devices placed on the EU market must meet the essential requirements, including the requirement that a medical device must be designed and manufactured in such a way that it will not compromise the clinical condition or safety of patients, or the safety and health of users and others. In addition, the device must achieve the performance intended by the manufacturer and be designed, manufactured, and packaged in a suitable manner.

Compliance with the essential requirements is a prerequisite for European Conformity Marking ("CE mark") without which medical devices cannot be marketed or sold in the EU. To demonstrate compliance with the essential requirements, medical device manufacturers must undergo a conformity assessment procedure, which

varies according to the type of medical device and its (risk) classification. Except for low-risk medical devices (Class I non-sterile, non-measuring devices), where the manufacturer can self-assess the conformity of its products with the essential requirements (except for any parts which relate to sterility or metrology), a conformity assessment procedure requires the intervention of a notified body. Notified bodies are independent organizations designated by EU member states to assess the conformity of devices before being placed on the market. A notified body would typically audit and examine a product's technical dossiers and the manufacturers' quality system. If satisfied that the relevant product conforms to the relevant essential requirements, the notified body issues a certificate of conformity, which the manufacturer uses as a basis for its own declaration of conformity. The manufacturer may then apply the CE mark to the device, which allows the device to be placed on the market throughout the EU.

Throughout the term of the certificate of conformity, the manufacturer will be subject to periodic surveillance audits to verify continued compliance with the applicable requirements. In particular, there will be a new audit by the notified body before it will renew the relevant certificate(s).

### *Medical Devices Regulation*

On April 5, 2017, the EU Medical Devices Regulation was adopted with the aim of ensuring better protection of public health and patient safety. The EU Medical Devices Regulation establishes a uniform, transparent, predictable and sustainable regulatory framework across the EU for medical devices and ensures a high level of safety and health while supporting innovation. Unlike the EU Medical Devices Directive and the AIMDD, the EU Medical Devices Regulation is directly applicable in EU member states without the need for member states to implement into national law. This aims at increasing harmonization across the EU.

The EU Medical Devices Regulation became effective on May 26, 2021. The new Regulation among other things:

- strengthens the rules on placing devices on the market (e.g. reclassification of certain devices and wider scope than the EU Medical Devices Directive and the AIMDD) and reinforces surveillance once they are available;
- establishes explicit provisions on manufacturers' responsibilities for the follow-up of the quality, performance and safety of devices placed on the market;
- establishes explicit provisions on importers' and distributors' obligations and responsibilities;
- imposes an obligation to identify a responsible person who is ultimately responsible for all aspects of compliance with the requirements of the new regulation;
- improves the traceability of medical devices throughout the supply chain to the end-user or patient through the introduction of a unique identification number, to increase the ability of manufacturers and regulatory authorities to trace specific devices through the supply chain and to facilitate the prompt and efficient recall of medical devices that have been found to present a safety risk;
- sets up a central database (Eudamed) to provide patients, healthcare professionals and the public with comprehensive information on products available in the EU; and
- strengthens rules for the assessment of certain high-risk devices, such as implants, which may have to undergo a clinical evaluation consultation procedure by experts before they are placed on the market.

Devices lawfully placed on the market pursuant to the EU Medical Devices Directive or the AIMDD prior to May 26, 2021 may generally continue to be made available on the market or put into service until May 26, 2025, provided that the requirements of the transitional provisions are fulfilled. In particular, the certificate in question must still be valid. However, even in this case, manufacturers must comply with a number of new or reinforced requirements set forth in the EU Medical Devices Regulation, in particular the obligations described below.

The EU Medical Devices Regulation requires that before placing a device, other than a custom-made device, on the market, manufacturers (as well as other economic operators such as authorized representatives and importers) must register by submitting identification information to the electronic system (Eudamed), unless they have already registered. The information to be submitted by manufacturers (and authorized representatives) also includes the name, address and contact details of the person or persons responsible for regulatory compliance. The new Regulation also requires that before placing a device, other than a custom-made device, on the market, manufacturers must assign a unique identifier to the device and provide it along with other core data to the unique device identifier (“UDI”) database. These new requirements aim at ensuring better identification and traceability of the devices. Each device – and as applicable, each package – will have a UDI composed of two parts: a device identifier (“UDI-DI”) specific to a device, and a production identifier (“UDI-PI”) to identify the unit producing the device. Manufacturers are also notably responsible for entering the necessary data on Eudamed, which includes the UDI database, and for keeping it up to date. The obligations for registration in Eudamed will become applicable at a later date (as Eudamed is not yet fully functional). Until Eudamed is fully functional, the corresponding provisions of the EU Medical Devices Directive and the AIMDD continue to apply for the purpose of meeting the obligations laid down in the provisions regarding exchange of information, including, and in particular, information regarding registration of devices and economic operators.

All manufacturers placing medical devices on the market in the EU must comply with the EU medical device vigilance system which has been reinforced by the EU Medical Devices Regulation. Under this system, serious incidents and Field Safety Corrective Actions (“FSCAs”) must be reported to the relevant authorities of the EU member states. These reports will have to be submitted through Eudamed – once functional – and aim to ensure that, in addition to reporting to the relevant authorities of the EU member states, other actors such as the economic operators in the supply chain will also be informed. Until Eudamed is fully functional, the corresponding provisions of the EU Medical Devices Directive and the AIMDD continue to apply. A serious incident is defined as any malfunction or deterioration in the characteristics or performance of a device made available on the market, including use-error due to ergonomic features, as well as any inadequacy in the information supplied by the manufacturer and any undesirable side-effect, which, directly or indirectly, might have led or might lead to the death of a patient or user or of other persons or to a temporary or permanent serious deterioration of a patient's, user's or other person's state of health or a serious public health threat. Manufacturers are required to take FSCAs defined as any corrective action for technical or medical reasons to prevent or reduce a risk of a serious incident associated with the use of a medical device that is made available on the market. An FSCA may include the recall, modification, exchange, destruction or retrofitting of the device. FSCAs must be communicated by the manufacturer or its legal representative to its customers and/or to the end users of the device through Field Safety Notices. For similar serious incidents that occur with the same device or device type and for which the root cause has been identified or a FSCA implemented or where the incidents are common and well documented, manufacturers may provide periodic summary reports instead of individual serious incident reports.

The aforementioned EU rules are generally applicable in the European Economic Area (“EEA”) which consists of the 27 EU member states plus Norway, Liechtenstein, and Iceland.

### *Brexit*

Since January 1, 2021, the Medicines and Healthcare Products Regulatory Agency (“MHRA”) has become the sovereign regulatory authority responsible for Great Britain (i.e. England, Wales and Scotland) medical device market according to the requirements provided in the Medical Devices Regulations 2002 (SI 2002 No 618, as amended) that sought to give effect to the three pre-existing EU directives governing active implantable medical devices, general medical devices and in vitro diagnostic medical devices whereas Northern Ireland continues to be governed by EU rules according to the Northern Ireland Protocol. Following the end of the Brexit transitional period on January 1, 2021, new regulations require medical devices to be registered with the MHRA (but manufacturers were given a grace period of four to 12 months, depending on the classification of the device, to comply with the new registration process) before being placed on Great Britain market. The MHRA only registers devices where the manufacturer or their United Kingdom (“UK”) Responsible Person has a registered place of business in the UK. Manufacturers based outside the UK need to appoint a UK Responsible Person that has a registered place of business in the UK to register devices with the MHRA in line with the grace periods. By July 1, 2023, in Great Britain, all medical devices will require a UKCA (“UK Conformity Assessed”) mark but CE marks issued by EU notified bodies will remain valid until this time. Manufacturers may choose to use the UKCA mark on

a voluntary basis until June 30, 2023. However, UKCA marking will not be recognized in the EU. The rules for placing medical devices on the market in Northern Ireland, which is part of the UK, differ from those in the rest of the UK. Compliance with this legislation is a prerequisite to be able to affix the UKCA mark to our products, without which they cannot be sold or marketed in Great Britain.

An MHRA public consultation was opened until end of November 2021 on the post-Brexit regulatory framework for medical devices and diagnostics. MHRA seeks to amend the UK Medical Devices Regulations 2002 (which are based on EU legislation, primarily the EU Medical Devices Directive and the EU In Vitro Diagnostic Medical Devices Directive 98/79/EC), in particular to create new access pathways to support innovation, create an innovative framework for regulating software and artificial intelligence as medical devices, reform IVD regulation, and foster sustainability through the reuse and remanufacture of medical devices. The regime is expected to come into force in July 2023, coinciding with the end of the acceptance period for EU CE marks in Great Britain, subject to appropriate transitional arrangements. The consultation indicated that the MHRA will publish guidance in relation to the changes to the regulatory framework and may rely more heavily on guidance to add flexibility to the regime.

In addition, the Trade Deal between the UK and the EU generally provides for cooperation and exchange of information between the parties in the areas of product safety and compliance, including market surveillance, enforcement activities and measures, standardization-related activities, exchanges of officials, and coordinated product recalls. As such, processes for compliance and reporting should reflect requirements from regulatory authorities.

Similarly, we are subject to regulations and product registration requirements in many foreign countries in which we may sell our products, including in the areas of:

- design, development, manufacturing, and testing;
- product standards;
- product safety;
- product safety reporting;
- marketing, sales, and distribution;
- packaging and storage requirements;
- labeling requirements;
- content and language of instructions for use;
- clinical trials;
- record keeping procedures;
- advertising and promotion;
- recalls and field corrective actions;
- post-market surveillance, including reporting of deaths or serious injuries and malfunctions that, if they were to recur, could lead to death or serious injury;
- import and export restrictions;
- tariff regulations, duties, and tax requirements;

- registration for reimbursement; and
- necessity of testing performed in country by distributors for licensees.

The time required to obtain clearance or certification required by foreign countries may be longer or shorter than that required for FDA clearance, and requirements for licensing a product in a foreign country may differ significantly from FDA requirements.

### ***Federal, State, and Foreign Fraud and Abuse Laws***

In addition to FDA restrictions on marketing and promotion of drugs and devices, other federal, state, and foreign laws restrict our business practices. These laws include, without limitation, foreign, federal, and state anti-kickback and false claims laws, as well as transparency laws regarding payments or other items of value provided to healthcare providers.

The federal Anti-Kickback Statute prohibits, among other things, knowingly and willfully offering, paying, soliciting or receiving any remuneration (including any kickback, bribe or rebate), directly or indirectly, overtly or covertly, in cash or in kind to induce or in return for purchasing, leasing, ordering or arranging for or recommending the purchase, lease or order of any good, facility, item or service reimbursable, in whole or in part, under Medicare, Medicaid or other federal healthcare programs. The term “remuneration” has been broadly interpreted to include anything of value, including stock, stock options, and the compensation derived through ownership interests.

Although there are a number of statutory exceptions and regulatory safe harbors protecting some common activities from prosecution, the exceptions and safe harbors are drawn narrowly. Failure to meet all of the requirements of a particular applicable statutory exception or regulatory safe harbor does not make the conduct per se illegal under the federal Anti-Kickback Statute. Instead, the legality of the arrangement will be evaluated on a case-by-case basis based on a cumulative review of all its facts and circumstances. Conduct and business arrangements that do not fully satisfy one of these safe harbor provisions may result in increased scrutiny by government enforcement authorities. Several courts have interpreted the statute’s intent requirement to mean that if any one purpose of an arrangement involving remuneration is to induce referrals of federal healthcare covered business, the federal Anti-Kickback Statute has been violated. In addition, a person or entity does not need to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation.

The majority of states also have anti-kickback laws which establish similar prohibitions and in some cases may apply more broadly to items or services covered by any third-party payor, including commercial insurers and self-pay patients.

The federal civil False Claims Act prohibits, among other things, any person or entity from knowingly presenting, or causing to be presented, a false or fraudulent claim for payment or approval to the federal government or knowingly making, using or causing to be made or used a false record or statement material to a false or fraudulent claim to the federal government. A claim includes “any request or demand” for money or property presented to the U.S. government. The federal civil False Claims Act also applies to false submissions that cause the government to be paid less than the amount to which it is entitled, such as a rebate. Intent to deceive is not required to establish liability under the civil federal civil False Claims Act. Moreover, 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 civil False Claims Act.

In addition, private parties may initiate “qui tam” whistleblower lawsuits against any person or entity under the federal civil False Claims Act in the name of the government and share in the proceeds of the lawsuit. The government may further prosecute conduct constituting a false claim under the federal criminal False Claims Act. The criminal False Claims Act prohibits the making or presenting of a claim to the government knowing such claim to be false, fictitious or fraudulent and, unlike the federal civil False Claims Act, requires proof of intent to submit a false claim.

The Civil Monetary Penalty Law imposes penalties against any person or entity that, among other things, is determined to have presented or caused to be presented a claim 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, or offering or transferring remuneration to a federal healthcare beneficiary that a person knows or should know is likely to influence the beneficiary's decision to order or receive items or services reimbursable by the government from a particular provider or supplier.

The Health Insurance Portability and Accountability Act of 1996 ("HIPAA") also created additional federal criminal statutes that prohibit among other actions, knowingly and willfully executing, or attempting to execute, a scheme to defraud any healthcare benefit program, including private third-party payors, knowingly and willfully embezzling or stealing from a healthcare benefit program, willfully obstructing a criminal investigation of a healthcare offense, and 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. Similar to the federal Anti-Kickback Statute, a person or entity does not need to have actual knowledge of the statute or specific intent to violate it in order to have committed a violation.

Many foreign countries have similar laws relating to healthcare fraud and abuse. Foreign laws and regulations may vary greatly from country to country. For example, the advertising and promotion of medical devices is subject to some general principles set forth in EU legislation. According to the EU Medical Devices Regulation, only devices that are CE marked may be marketed and advertised in the EU in accordance with their intended purpose. Directive 2006/114/EC concerning misleading and comparative advertising and Directive 2005/29/EC on unfair commercial practices, while not specific to the advertising of medical devices, also apply to the advertising thereof and contain general rules, for example, requiring that advertisements are evidenced, balanced and not misleading. Specific requirements are defined at a national level. EU member states' laws related to the advertising and promotion of medical devices, which vary between jurisdictions, may limit or restrict the advertising and promotion of products to the general public and may impose limitations on promotional activities with healthcare professionals. These laws, which vary between jurisdictions (thus making compliance more complex), may limit or restrict the advertising and promotion of our products to the general public and may impose limitations on our promotional activities with healthcare professionals. Many EU member states have adopted specific anti-gift statutes that further limit commercial practices for our products, in particular vis-à-vis healthcare professionals and organizations. Additionally, there has been a recent trend of increased regulation of payments and transfers of value provided to healthcare professionals or entities and many EU member states have adopted national "Sunshine Acts" which impose reporting and transparency requirements (often on an annual basis), similar to the requirements in the United States, on medical device manufacturers. Certain countries also mandate implementation of commercial compliance programs. Also, many U.S. states have similar fraud and abuse statutes or regulations that may be broader in scope and may apply regardless of payor, in addition to items and services reimbursed under Medicaid and other state programs.

Additionally, there has been a recent trend of increased foreign, federal, and state regulation of payments and transfers of value provided to healthcare professionals or entities. In the U.S., the federal Physician Payments Sunshine Act imposes annual reporting requirements on certain drug, biologics, medical supplies and device manufacturers for which payment is available under Medicare, Medicaid or CHIP for payments and other transfers of value provided by them, directly or indirectly, to physicians, as defined by statute, certain other non-physician practitioners such as physician assistants and nurse practitioners, and teaching hospitals, as well as ownership and investment interests held by physicians and their immediate family members. Manufacturers must submit reports by the 90th day of each calendar year. Many EU member states have adopted national "Sunshine Acts" which impose similar reporting and transparency requirements (often on an annual basis) on certain drug, biologics and medical device manufacturers. Certain foreign countries and U.S. states also mandate implementation of commercial compliance programs, impose restrictions on device manufacturer marketing practices and require tracking and reporting of gifts, compensation, and other remuneration to healthcare professionals and entities.

Violation of any of the federal and state healthcare laws described above or any other governmental regulations that apply to device manufacturers may result in significant penalties, including the imposition of significant civil, criminal and administrative penalties, damages, disgorgement, monetary fines, imprisonment, possible exclusion from participation in Medicare, Medicaid and other federal healthcare programs, contractual damages, reputational harm, diminished profits and future earnings, additional reporting requirements and/or oversight if the



entity becomes subject to a corporate integrity agreement or similar agreement to resolve allegations of non-compliance with these laws, and curtailment of operations.

### ***Data Privacy and Security Laws***

Numerous state, federal and foreign laws, regulations, and standards govern the collection, use, access to, confidentiality and security of health-related and other personal information, and could apply now or in the future to our operations or the operations of our partners. In the United States, numerous federal and state laws and regulations, including data breach notification laws, health information privacy and security laws, including HIPAA, and consumer protection laws and regulations govern the collection, use, disclosure, and protection of health-related and other personal information. In addition, certain foreign laws govern the privacy and security of personal data, including health-related data. For example, the General Data Protection Regulation (the "GDPR"), imposes strict requirements for processing the personal data of individuals within the European Economic Area, or the EEA. Companies that must comply with the GDPR face increased compliance obligations and risk, including more robust regulatory enforcement of data protection requirements and potential fines for noncompliance of up to €20 million or 4% of the annual global revenues of the noncompliant company, whichever is greater. Further, from January 1, 2021, companies have had to comply with the GDPR and also the UK GDPR, which, together with the amended UK Data Protection Act 2018, retains the GDPR in UK national law. The UK GDPR mirrors the fines under the GDPR, i.e., fines up to the greater of €20 million (£17.5 million) or 4% of global turnover. Privacy and security laws, regulations, and other obligations are constantly evolving, may conflict with each other to complicate compliance efforts, and can result in investigations, proceedings, or actions that lead to significant civil and/or criminal penalties and restrictions on data processing.

### ***Healthcare Reform***

The U.S. and some foreign jurisdictions are considering or have enacted a number of legislative and regulatory proposals to change the healthcare system in ways that could affect our ability to sell our products profitably. Among policy makers and payors in the U.S. and elsewhere, there is significant interest in promoting changes in healthcare systems with the stated goals of containing healthcare costs, improving quality or expanding access. Current and future legislative proposals to further reform healthcare or reduce healthcare costs may limit coverage of or lower reimbursement for the procedures associated with the use of our products. The cost containment measures that payors and providers are instituting and the effect of any healthcare reform initiative implemented in the future could impact our revenue from the sale of our products.

The implementation of the Affordable Care Act in the U.S., for example, has changed healthcare financing and delivery by both governmental and private insurers substantially, and affected medical device manufacturers significantly. The Affordable Care Act, among other things, provided incentives to programs that increase the federal government's comparative effectiveness research, and implemented payment system reforms including a national pilot program on payment bundling to encourage hospitals, physicians, and other providers to improve the coordination, quality, and efficiency of certain healthcare services through bundled payment models. Additionally, the Affordable Care Act expanded eligibility criteria for Medicaid programs and created a new Patient-Centered Outcomes Research Institute to oversee, identify priorities in, and conduct comparative clinical effectiveness research, along with funding for such research.

Since its enactment, there have been judicial, executive and Congressional challenges to certain aspects of the Affordable Care Act. On June 17, 2021, the U.S. Supreme Court dismissed the most recent judicial challenge to the Affordable Care Act without specifically ruling on the constitutionality of the Affordable Care Act. Prior to the Supreme Court's decision, President Biden issued an executive order to initiate a special enrollment period from February 15, 2021 through August 15, 2021 for purposes of obtaining health insurance coverage through the Affordable Care Act marketplace. The executive order also instructed certain governmental agencies to review and reconsider their existing policies and rules that limit access to healthcare, including among others, reexamining Medicaid demonstration projects and waiver programs that include work requirements, and policies that create unnecessary barriers to obtaining access to health insurance coverage through Medicaid or the Affordable Care Act.

In addition, other legislative changes have been proposed and adopted since the Affordable Care Act was enacted. For example, the Budget Control Act of 2011, among other things, included reductions to Medicare payments to providers of 2% per fiscal year, which went into effect on April 1, 2013 and, due to subsequent legislative amendments to the statute, will remain in effect through 2030, with the exception of a temporary suspension from May 1, 2020 through March 31, 2022, unless additional Congressional action is taken. Additionally, the American Taxpayer Relief Act of 2012, among other things, reduced Medicare payments to several providers, including hospitals, and increased the statute of limitations period for the government to recover overpayments to providers from three to five years.

We expect additional state and federal healthcare reform measures to be adopted in the future, any of which could limit the amounts that federal and state governments will pay for healthcare products and services, which could result in reduced demand for our products or additional pricing pressure.

### ***Anti-Bribery and Corruption Laws***

Our U.S. operations are subject to the FCPA. We are required to comply with the FCPA, which generally prohibits covered entities and their intermediaries from engaging in bribery or making other prohibited payments to foreign officials for the purpose of obtaining or retaining business or other benefits. In addition, the FCPA imposes accounting standards and requirements on publicly traded U.S. corporations and their foreign affiliates, which are intended to prevent the diversion of corporate funds to the payment of bribes and other improper payments, and to prevent the establishment of “off books” slush funds from which such improper payments can be made. We also are subject to similar anticorruption legislation implemented in Europe through EU member state laws and under the Organization for Economic Co-operation and Development’s Convention on Combating Bribery of Foreign Public Officials in International Business Transactions.

### **Segment Information**

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

### **Seasonality**

Historically, we have experienced seasonality in our first and fourth quarters, and we expect this trend to continue. In the U.S., we have experienced, and may in the future experience, higher sales in the fourth quarter as a result of patients having paid their annual insurance deductibles in full, thereby reducing their out-of-pocket costs. In the first quarter of each year in Germany, we have experienced reduced demand for our Inspire therapy as Neue Untersuchungs-und-Behandlungsmethoden (“NUB”) coverage status is being determined and as hospitals are establishing their budgets pertaining to allocation of funds to purchase our Inspire therapy. Beginning January 1, 2021, Inspire therapy is fully integrated into the German hospital reimbursement system (“G-DRG”), and we therefore may experience less seasonal fluctuations in Germany although it may not eliminate them.

### **Human Capital**

#### ***Employees***

As of December 31, 2021, we had 485 employees, of which 459 are in the U.S., 25 are in Europe and one is in Japan. We increased the number of employees by 42% during 2021 to support the rapid growth of our business. Our internal promotion and progression rate was 15%, of which 60% were women and diverse employees. We continue to experience a minimal voluntary attrition rate of 6%, despite the market’s strong competition for talent.

None of our employees is subject to a collective bargaining agreement or represented by a trade or labor union. We consider our relationship with our employees to be good.

### *Code of Business Conduct and Business Ethics*

All employees are expected to conduct business with the highest standards business ethics. Each employee receives and agrees to follow the Inspire Medical Systems Code of Business Conduct and Ethics. Employees are encouraged to discuss any related concerns with management or report concerns anonymously through an Ethics Hotline. All reports received on the Ethics Hotline are investigated by an independent third party.

### *Culture of Respect and Inclusion*

We strive to create a culture in which all employees feel heard, respected, and valued. All new employees participate in training focused on appropriate, respectful, and inclusive workplace behavior. In 2020, we created an engaging training initiative for all employees which encourages awareness of unconscious bias and microaggressions. Building on our 2020 program, we implemented sessions specifically for Inspire leaders in their roles to create respectful culture within their teams. The goals of these programs are to encourage broad and diverse viewpoints to achieve the best outcomes for our patients, healthcare providers, and employees, and to build awareness of how our own behaviors impact our colleagues.

### *Talent Management & Development*

With our aggressive growth objectives, it is imperative that we have a strong pipeline of external candidates as well as opportunities for current employees to grow within our organization. To support these critical success factors, we have continued to invest in our talent acquisition group. In addition, we have created an internal Learning and Development organization to lead our development initiatives. To encourage further professional development of our employees, in 2022 we introduced a tuition reimbursement program for those pursuing a degree.

Through internal promotions, we proactively reorganized our U.S. sales team structure to execute continued rapid growth. To help support the expanded role of our leadership, we created a new leadership program specially designed to help our new leaders be successful in their expanded roles, as well as provide coaching opportunities for our leaders through external partners.

On an annual basis, our leadership team participates in a talent review and succession planning exercise to identify organizational needs, development opportunities, and potential future leaders. This enables us to identify the resources and skill sets needed to meet our growth objectives.

### *Compensation Philosophy to Drive High Performing Teams*

Our human capital strategies, initiatives, and outcomes are reviewed on a regular basis with our board of directors ensuring alignment with the Company's overall business strategies. We continue to evaluate all reward-related strategies to remain competitive externally and internally and ensure our rewards are aligned with our compensation philosophy. At the direction of the compensation committee of our board of directors, we continue to partner with a consulting firm to benchmark our peer group companies and other key companies from which we recruit talent. As a result of this analysis, we have implemented a number of changes to our rewards practices which allows us to maintain our competitiveness in the market.

We continue to believe strongly in providing employees the opportunity to participate as owners in the Company. This is done through equity programs such as the employee stock purchase program and the grant of stock options and restricted stock units. Additionally, we have incorporated a 401(k) employer match for all U.S.-based employees starting in 2022.

### *COVID-19 Health and Safety*

COVID-19 continues to impact our employees, healthcare providers, and patients, which requires us to find creative and flexible strategies to address the effects. Our top priority is the health of our employees and their families while ensuring business continuity. Our safety practices align closely to the guidance set by the Centers for Disease Control and locally mandated regulations. We have continued with contact tracing of all potential or

confirmed COVID-19 cases throughout our organization to minimize exposure to our employees as well as our external partners, and we have implemented a flexible work schedule to accommodate quarantine situations and the care of family members impacted by COVID-19.

### *Community Engagement*

Throughout the year, Inspire engages in community programs which align with our Company values. Traditional means of community involvement, however, continue to be a challenge through the pandemic. Where feasible, we encourage employees to participate in local level in community service programs, including the American Heart Association's Twin Cities Heart Walk, and also companywide participation in the annual Toys for Tots campaign.

### **Available Information**

We file Annual Reports on Form 10-K, Quarterly Reports on Form 10-Q, Current Reports on Form 8-K, proxy statements, amendments to such documents and other information with the SEC. Our SEC filings are available to the public over the Internet at the SEC's website at <http://www.sec.gov>. We also make these filings available, free of charge, under the Investor Relations section of our website at [www.inspiresleep.com](http://www.inspiresleep.com) as soon as reasonably practicable after we electronically file such material with, or furnish such material to, the SEC. Our website and the information contained on or connected to that site are not incorporated into this Annual Report on Form 10-K or any other public filing made by us with the SEC.

### **Item 1A. Risk Factors.**

*Investing in our common stock involves a high degree of risk. These risks include, but are not limited to, those described below, each of which may be relevant to an investment decision. You should carefully consider the risks described below, together with the other information included or incorporated by reference in this Annual Report on Form 10-K. The realization of any of the following risks could have a significant adverse effect on our reputation, business, financial condition, results of operations, growth, and our ability to accomplish our strategic objectives. In that event, the trading price of our common stock could decline. Other events that we do not currently anticipate or that we currently deem immaterial may also affect our reputation, business, financial conditions, results of operations, growth, and our ability to accomplish our strategic objectives.*

### **Risks Related to Our Business**

***We have incurred significant operating losses since inception, we expect to incur operating losses in the future and we may not be able to achieve or sustain profitability. We have limited history operating as a commercial company.***

We have incurred net losses since our inception in 2007. For the years ended December 31, 2021, 2020, and 2019, we had net losses of \$42.0 million, \$57.2 million, and \$33.2 million, respectively. As of December 31, 2021, we had an accumulated deficit of \$279.4 million. To date, we have financed our operations primarily through sales of our Inspire system, private placements of our convertible preferred securities, amounts borrowed under our credit facility, the initial public offering of our common stock that closed in May 2018 ("IPO"), and the two follow-on offerings of our common stock that closed in December 2018 and April 2020 (our "follow-on offerings"). We have devoted significant resources to research and development activities related to our Inspire system, including clinical and regulatory initiatives to obtain marketing approval, and sales and marketing activities.

We first commercialized our Inspire system in certain European markets in 2011, in the U.S. in 2014, and Japan in 2021 and therefore do not have a long history operating as a commercial company. Since 2011, our revenue has been derived, and we expect it to continue to be derived, primarily from sales of our Inspire system. Because of its recent commercial introduction, our Inspire system has limited product and brand recognition, particularly in new markets. In addition, demand for our Inspire system may decline or may not increase as quickly as we expect. Our ability to generate revenue from sales of our Inspire system, or from any products we may develop in the future, may not be sufficient to enable us to transition to profitability and generate positive cash flows.

We expect that our operating expenses will continue to increase as we continue to build our commercial infrastructure, invest in research and development, and develop, enhance, and commercialize new products. As a result, we expect to continue to incur operating losses for the foreseeable future and may never achieve profitability. Furthermore, even if we do achieve profitability, we may not be able to sustain or increase profitability on an ongoing basis. If we do not achieve or sustain profitability, it will be more difficult for us to finance our business and accomplish our strategic objectives, either of which would have a material adverse effect on our business, financial condition, results of operations, and cause the market price of our common stock to decline. In addition, failure of our Inspire system to significantly penetrate existing or new markets would negatively affect our business, financial condition, and results of operations.

***Our revenue is primarily generated from sales of our Inspire system and we are, therefore, highly dependent on it for our success.***

We began selling our Inspire system in 2011 in certain European countries, in 2014 in the U.S., and in 2021 in Japan. Sales of our Inspire system accounted for primarily all of our revenue for the years ended December 31, 2021, 2020, and 2019. We expect that sales of our Inspire system will continue to account for the substantial majority of our revenue going forward. Our ability to execute our growth strategy and become profitable will therefore depend upon the adoption by patients, physicians, and sleep centers, among others, of our Inspire therapy to treat moderate to severe OSA in patients who are unable to use or get consistent benefit from CPAP. Some physicians may have prior history with or a preference for other treatment options, such as positive airway pressure devices, surgical treatments or oral appliances, or may be reluctant to alter their practice patterns and undergo the training required to enable them to treat patients with our Inspire therapy. Patients may not adopt our Inspire therapy if, among other potential reasons, their airway anatomy would not allow for effective treatment with Inspire therapy, they are reluctant to receive an implantable device as opposed to an alternative, non-implantable treatment, they are worried about potential adverse effects of our Inspire system, such as infection, discomfort from the stimulation or tongue soreness or weakness, or they are unable to obtain adequate third-party coverage or reimbursement for our Inspire therapy.

We cannot ensure that our Inspire therapy will achieve broad market acceptance among physicians and patients. Any failure of the Inspire system to satisfy physician or patient demand or to achieve meaningful market acceptance will harm our business and future prospects.

***If patients or physicians are not willing to change current practices to adopt our Inspire therapy to treat moderate to severe OSA, our Inspire therapy may fail to gain increased market acceptance, and our business will be adversely affected.***

Our primary strategy to grow our revenue is to drive an increase in the adoption of our Inspire therapy to treat patients with moderate to severe OSA who are unable to use or get consistent benefit from CPAP. While the number of physicians prescribing our Inspire therapy has increased in recent years, there is a significant group of physicians who have not yet adopted our Inspire therapy, and additional physicians may choose not to adopt our Inspire therapy for a number of reasons, including:

- lack of availability of adequate third-party payor coverage or reimbursement;
- lack of experience with our products and with upper airway neurostimulation as a treatment alternative;
- our inability to convince key opinion leaders to provide recommendations regarding our Inspire therapy, or to convince physicians, patients, and healthcare payors that our Inspire therapy is an attractive alternative to other treatment options;
- perceived inadequacy of evidence supporting clinical benefits or cost-effectiveness of our Inspire therapy over existing alternatives;
- a perception among some physicians of patients' inability to tolerate the surgical procedure required to implant our Inspire system;

- liability risks generally associated with the use of new products and procedures; and
- the training required to use new products.

We focus our sales, marketing, and training efforts primarily on ENT physicians and sleep physicians. However, physicians from other disciplines, including cardiologists, pulmonologists, electrophysiologists, and primary care physicians, as well as other medical professionals, such as dentists, nurse practitioners, and physician assistants, are often the initial point of contact for patients with OSA.

These physicians and other medical professionals commonly screen and treat patients with moderate to severe OSA, and are likely to prescribe more conventional second-line treatment methods for patients who are unable to use or obtain consistent benefit from CPAP. We believe that educating physicians in these disciplines and other medical professionals about the clinical merits and patient benefits of our Inspire therapy as a treatment for moderate to severe OSA is a key element of increasing the adoption of our Inspire therapy. If additional physicians or other medical professionals do not adopt, or existing physician customers cease prescribing our Inspire therapy for any reason, including those listed above, our ability to execute our growth strategy will be impaired, and our business may be adversely affected.

In addition, patients may not be able to adopt or may choose not to adopt our Inspire therapy if, among other potential reasons, their airway anatomy would not allow for effective treatment with Inspire therapy, they are reluctant to receive an implantable device as opposed to an alternative, non-implantable treatment, they are worried about potential adverse effects of our Inspire system, such as infection, discomfort from the stimulation or tongue soreness or weakness, or they are unable to obtain adequate third-party coverage or reimbursement.

***If we are unable to achieve and maintain adequate levels of coverage or reimbursement for our Inspire system, or any future products we may seek to commercialize, our commercial success may be severely hindered.***

We currently derive primarily all of our revenue from sales of our Inspire system and expect this to continue for the foreseeable future. The primary customers for our products are hospitals and ASCs. Our customers typically bill various third-party payors to cover all or a portion of the costs and fees associated with the procedures in which our products are used and bill patients for any deductibles or co-payments. Because there is often no separate reimbursement for supplies used in surgical procedures, the additional cost associated with the use of our products can affect the profit margin of the hospital or surgery center where the procedure is performed. Some of our target customers may be unwilling to adopt our products in light of the additional associated cost. Further, any decline in the amount payors are willing to reimburse our customers could make it difficult for existing customers to continue using or to adopt our products and could create additional pricing pressure for us. If we are forced to lower the price we charge for our products, our gross margins will decrease, which could have a material adverse effect on our business, financial condition, and results of operations, and impair our ability to grow our business.

Several third-party payors do not currently cover our products and the related procedures because they have determined that our products and the related procedures are experimental or investigational. When our products and the related procedures are covered, they are reimbursed primarily on a per-patient prior authorization basis for patients covered by commercial insurers, under Local Coverage Determinations for patients covered by Medicare, and under U.S. government contract for patients who are treated by the Veterans Health Administration. Customers who perform the procedure may be subject to reimbursement claim denials upon submission of the claim. Customers may also be subject to recovery of overpayments if a payor makes payment for the claim and subsequently determines that the payor's coding, billing or coverage policies were not followed. Our customers typically must directly bill patients enrolled with these third-party payors for the costs and fees associated with the procedures in which our products are used.

Third-party payors, whether foreign or domestic, or governmental or commercial, are developing increasingly sophisticated methods of controlling healthcare costs. In addition, no uniform policy of coverage and reimbursement for procedures using our products exists among third-party payors. Therefore, coverage and reimbursement for procedures using our products can differ significantly from payor to payor. Payors continually review new and existing technologies for possible coverage and can, without notice, deny or reverse coverage for

new or existing products and procedures. There can be no assurance that third-party payor policies will provide coverage for procedures in which our products are used. If we are not successful in reversing existing non-coverage policies, or if third-party payors that currently cover or reimburse our products and related procedures reverse or limit their coverage in the future, or if other third-party payors issue similar policies, this could have a material adverse effect on our business.

Further, we believe that future coverage and reimbursement may be subject to increased restrictions, such as additional prior authorization requirements, both in the U.S. and in international markets. In Europe, reimbursement is entirely regulated at the Member State level, varies significantly between countries, and member states are facing increased pressure to limit public healthcare spending. Third-party coverage and reimbursement for procedures using our products or any of our products in development for which we may receive regulatory approval may not be available or adequate in either the U.S. or international markets, which could have an adverse effect on our business, financial condition and results of operations, and impair our ability to grow our business.

***Third-party payors who do not cover, or physicians who do not use, our Inspire system may require additional clinical data prior to adopting or maintaining coverage of our Inspire system.***

Our success depends on physician and third-party payor acceptance of our Inspire therapy as an effective treatment option for patients with moderate to severe OSA. If physicians or payors do not find our body of published clinical evidence and data compelling or wish to wait for additional studies, they may choose not to use or provide coverage and reimbursement for our products. Currently, there are several third-party payors that have determined upper airway neurostimulation to be experimental or investigational and therefore do not cover it at this time.

In addition, the long-term effects of upper airway neurostimulation with our Inspire system beyond 10 years are not yet known. Certain physicians, hospitals, ASCs, and payors may prefer to see longer-term safety and efficacy data than we have produced. We cannot provide assurance that any data that we or others may generate in the future will be consistent with that observed in our existing clinical studies.

***The training required for physicians to use our Inspire system could reduce the market acceptance of our products.***

As with any new method or technique, physicians must undergo a thorough training program before they are qualified to perform the surgery to implant our Inspire system. Physicians could experience difficulty with the technique necessary to successfully insert the device and may not achieve the technical competency necessary to complete the training program. Even after successfully completing the training program, physicians could still experience difficulty implanting our Inspire system and, as a result, limit its use significantly in their practice or cease utilizing it altogether.

In addition, we may experience difficulty growing the number of physicians who complete our training program if patient demand is low, if the length of time necessary to train each physician is longer than expected, if the capacity of our sales representatives to train physicians is less than expected or if we are unable to sufficiently grow our sales organization. All of these events would lead to fewer trained physicians qualified to implant our Inspire system, which could negatively affect our business, financial condition and results of operations, and impair our ability to grow our business.

***We currently compete and will in the future continue to compete against other companies, some of which have longer operating histories, more established products or greater resources than we do, which may prevent us from achieving increased market penetration and improved operating results.***

The medical technology industry is highly competitive, subject to change and significantly affected by new product introductions and other activities of industry participants. Our competitors have historically dedicated and will continue to dedicate significant resources to promoting their products or developing new products or methods to treat moderate to severe OSA. We consider our primary competition to be other neurostimulation technologies designed to treat OSA. Though we are currently the only such technology approved for commercialization in the

U.S. by the FDA, we currently compete outside the U.S. with LivaNova, which produces an open-loop neurostimulation device and is currently conducting clinical trials of its device in the U.S. We also compete outside the U.S. with Nyxoah, which markets a bilateral hypoglossal nerve stimulation device in certain countries outside the U.S., and is conducting its first pivotal trial as it seeks FDA approval in the U.S. We believe other emerging businesses are in the early stages of developing neurostimulation devices designed to treat OSA. In addition, we also compete, both within and outside of the U.S., with invasive surgical treatment options such as UPPP and MMA and, to a lesser extent, oral appliances, which are primarily used in the treatment of mild to moderate OSA.

In addition, our Inspire therapy is approved for use as a second-line therapy in the treatment of moderate to severe OSA in patients who cannot use or obtain consistent benefit from CPAP. If one or more CPAP device manufacturers successfully develop a CPAP device that is more effective, better tolerated or otherwise results in better compliance by patients, or if improvements in other second-line therapies make them more effective, cost effective, easier to use or otherwise more attractive than our Inspire therapy, sales of our Inspire system could be significantly and adversely affected, which could have a material adverse effect on our business and financial condition and results of operations. In addition, if other companies are successful in developing neurostimulation devices that are approved for a broader range of indications than our Inspire system, we will be at a further competitive disadvantage, which could also affect our business, financial condition and results of operations.

Many of the companies against which we compete may have competitive advantages with respect to primary competitive factors in the OSA treatment market, including:

- greater company, product, and brand recognition;
- superior product safety, reliability, and durability;
- better quality and larger volume of clinical data;
- more effective marketing to and education of patients, physicians, and sleep centers;
- greater product ease of use and patient comfort;
- more sales force experience and greater market access;
- better product support and service;
- more advanced technological innovation, product enhancements, and speed of innovation;
- more effective pricing and revenue strategies;
- lower procedure costs to patients;
- more effective reimbursement teams and strategies;
- dedicated practice development; and
- more effective clinical training teams.

Most of the other OSA treatments against which we compete have a greater penetration into the OSA treatment market. Oral appliances and other surgical treatments are better known to ENT physicians, sleep centers, and the other physicians on whom we rely for referrals.

We also compete with other medical technology companies to recruit and retain qualified sales, training, and other personnel, including members of our in-house prior authorization team.

In addition, though there are currently no pharmacologic therapies approved to treat OSA, we may in the future face competition from pharmaceutical companies that develop such therapies. We also expect to experience



increased competition in the future as other companies develop and commercialize competing neurostimulation devices. Any of these companies may also have the competitive advantages described above.

***Our business, financial condition, results of operations and growth could be significantly harmed by the effects of the COVID-19 pandemic.***

We are subject to risks related to public health crises such as the global pandemic associated with COVID-19. In 2020, a novel strain of coronavirus, SARS-CoV-2, and the resulting disease COVID-19, spread around the world and within the U.S. New variants of the virus have subsequently emerged, some of which have shown to be more contagious. The COVID-19 pandemic has negatively impacted our business, results of operations and financial condition by significantly decreasing and delaying the number of Inspire therapy procedures performed and patients screened for eligibility for Inspire therapy, and we expect the COVID-19 pandemic to continue to negatively impact our business, results of operations and financial condition. The number of Inspire therapy procedures performed, similar to other elective surgical procedures, decreased compared to pre-pandemic levels as health care organizations in the U.S. and globally have prioritized the treatment of patients with COVID-19. For example, substantially all of our Inspire therapy procedures were postponed beginning in the second week of March 2020 and numerous other cases, which received prior authorization approval, were not able to be scheduled and therefore were also postponed. Beginning in May 2020, surgical volumes began to increase steadily through the remainder of 2020, with most implanting centers performing procedures by October 2020. Resurgences of COVID-19 in various U.S. and European regions and Japan have, and may in the future, negatively impact our procedure volumes. Additionally, while procedural volume rates for elective surgeries generally returned to pre-pandemic levels in the U.S. and Europe by March 31, 2021, the COVID-19 pandemic continues to rapidly evolve and its impact on our business will depend on several factors that are highly uncertain and unpredictable. The COVID-19 pandemic will likely continue to adversely impact our procedure volumes. These measures and challenges will likely continue for the duration of the pandemic, which is uncertain, and will continue to significantly reduce our revenue and negatively impact our business, results of operations and financial condition while the pandemic continues. Further, if there are additional or reinstated government mandated or recommended cancellations of elective surgical procedures, this could create in the future a substantial backlog of patients seeking appointments with physicians and surgeries to be performed at hospitals and ambulatory surgery centers relating to a variety of medical conditions, and as a result, patients seeking Inspire therapy procedures to be performed will have to navigate limited provider capacity. We believe this limited provider, hospital and ambulatory surgery center capacity could have a significant adverse effect on our business, results of operations and financial condition.

Numerous foreign, state and local jurisdictions have imposed, and others in the future may impose, “shelter-in-place” orders, quarantines, social distancing requirements, executive orders and similar government orders and restrictions for their residents to control the spread of COVID-19. Although such orders have been lifted, various disruptions remain ongoing. These disruptions have included, and future disruptions may also include, restrictions on our personnel and personnel of partners to travel and access customers for training and case support; inability of our suppliers to manufacture and test our Inspire therapy and its components and to deliver these on a timely basis, or at all; inventory shortages or obsolescence; delays in approvals or certifications by regulatory authorities or notified bodies; delays in ongoing preclinical trials; delays in operations at insurance agencies, which may impact timelines for the issuance of insurance coverage policies; diversion of or limitations on employee resources that would otherwise be focused on the operations of our business, including because of sickness of employees or their families or the desire of employees to avoid contact with large groups of people; delays in growing or reductions in our sales team, including through delays in hiring, lay-offs, furloughs or other losses of sales representatives; and additional government requirements or other incremental mitigation efforts that may further impact our or our suppliers’ capacity to manufacture our Inspire system. The extent to which the COVID-19 pandemic impacts our business will depend on future developments, which are highly uncertain and cannot be predicted, including new information which may emerge concerning the severity and spread of COVID-19, future waves of infection, and the actions to contain COVID-19 or treat its impact, among others.

While the potential economic impact brought by and the duration of COVID-19 may be difficult to assess or predict, the widespread pandemic has resulted in, and may continue to result in, significant disruption of global financial markets, reducing our ability to access capital, which could in the future negatively affect our liquidity. In addition, a recession or market correction resulting from the COVID-19 pandemic could materially affect our

business. The occurrence of any such events may lead to reduced disposable income and access to health insurance which could adversely affect the number of Inspire systems sold after the pandemic has ended.

***Our long-term growth depends on our ability to enhance our Inspire system, expand our indications, and develop and commercialize additional products.***

It is important to our business that we continue to enhance our Inspire system and develop and introduce new products. Developing products is expensive and time-consuming and could divert management's attention away from our core business. The success of any new product offering or product enhancements to our Inspire system will depend on several factors, including our ability to:

- properly identify and anticipate physician and patient needs;
- develop and introduce new products and product enhancements in a timely manner;
- avoid infringing upon the intellectual property rights of third-parties;
- demonstrate, if required, the safety and efficacy of new products with data from preclinical studies and clinical trials;
- obtain the necessary regulatory clearances, approvals or certifications for expanded indications, new products or product modifications;
- be fully FDA-compliant with marketing of new devices or modified products and be fully compliant with foreign requirements to market our new devices or modified products;
- provide adequate training to potential users of our products;
- receive adequate coverage and reimbursement for procedures performed with our products; and
- develop an effective and dedicated sales and marketing team.

If we are not successful in expanding our indications and developing and commercializing new products and product enhancements, our ability to increase our revenue may be impaired, which could have a material adverse effect on our business, financial condition, and results of operations.

***Our financial results may fluctuate significantly and may not fully reflect the underlying performance of our business.***

Our quarterly and annual results of operations may vary significantly in the future, and period-to-period comparisons of our operating results may not be meaningful. Accordingly, the results of any one quarter or period should not be relied upon as an indication of future performance. Our quarterly and annual financial results may fluctuate as a result of a variety of factors, many of which are outside our control and, as a result, may not fully reflect the underlying performance of our business. Such factors may include, for example, seasonal variations in our sales or required postponements of elective surgical procedures effected during a health crisis, as was the case with the COVID-19 pandemic. We generally experience and may in the future experience higher sales in the U.S. during the fourth quarter as a result of patients having paid their annual insurance deductibles in full, thereby reducing their out-of-pocket costs. Additionally, in the first quarter of each year in Europe, we have historically experienced reduced demand for our Inspire therapy as NUB coverage status is being determined in Germany and as hospitals are establishing their budgets pertaining to allocation of funds to purchase our Inspire therapy. Beginning January 1, 2021, Inspire therapy was fully integrated into the German hospital reimbursement system ("G-DRG"), however this may not eliminate our seasonal fluctuations in Germany.

Other factors that may cause fluctuations in our quarterly and annual results include:

- patient and physician adoption of our Inspire therapy;

- changes in coverage policies by third-party payors that affect the reimbursement of procedures using our products;
- timing of new product offerings, acquisitions, licenses or other significant events by us or our competitors;
- unanticipated pricing pressure;
- the hiring, retention, and continued productivity of our sales representatives;
- our ability to expand the geographic reach of our sales and marketing efforts;
- our ability to obtain regulatory clearance, approval, or certification for any products in development or for our current products for additional indications or in additional countries outside the U.S.;
- results of clinical research and trials on our existing products and products in development;
- delays in receipt of anticipated purchase orders;
- delays in, or failure of, component and raw material deliveries by our suppliers; and
- positive or negative coverage in the media or clinical publications of our products or products of our competitors or our industry.

Because our quarterly and annual results may fluctuate, period-to-period comparisons may not be the best indication of the underlying results of our business and should only be relied upon as one factor in determining how our business is performing. These fluctuations may also increase the likelihood that we will not meet our forecasted performance, which could negatively affect the market price for our common stock.

***Our results of operations could be materially harmed if we are unable to accurately forecast customer demand for our Inspire system and manage our inventory.***

To ensure adequate inventory supply, we must forecast inventory needs and place orders with our suppliers based on our estimates of future demand for our Inspire system. Our ability to accurately forecast demand for our Inspire system could be negatively affected by many factors, including our failure to accurately manage our expansion strategy, product introductions by competitors, an increase or decrease in customer demand for our Inspire system or for products of our competitors, our failure to accurately forecast customer acceptance of new products, unanticipated changes in general market conditions or regulatory matters, and weakening of economic conditions or consumer confidence in future economic conditions. Inventory levels in excess of customer demand may result in inventory write-downs or write-offs, which would cause our gross margin to be adversely affected and could impair the strength of our brand. Conversely, if we underestimate customer demand for our Inspire system, our third-party contract manufacturers may not be able to deliver products to meet our requirements, and this could result in damage to our reputation and customer relationships. In addition, if we experience a significant increase in demand, additional supplies of raw materials or additional manufacturing capacity may not be available when required on terms that are acceptable to us, or at all, or suppliers or our third-party manufacturers may not be able to allocate sufficient capacity in order to meet our increased requirements, which could have an adverse effect on our ability to meet customer demand for our Inspire system and our results of operations.

We seek to maintain sufficient levels of inventory in order to protect ourselves from supply interruptions. As a result, we are subject to the risk that a portion of our inventory will become obsolete or expire, which could have a material adverse effect on our earnings and cash flows due to the resulting costs associated with the inventory impairment charges and costs required to replace such inventory.

***We rely on a limited number of third-party suppliers and contract manufacturers for the manufacture and assembly of our products, and a loss or degradation in performance of these suppliers and contract manufacturers could have a material adverse effect on our business, financial condition, and results of operations.***

We rely on third-party suppliers and contract manufacturers for the raw materials and components used in our Inspire system and to manufacture and assemble our products. The suppliers that provide certain materials and components are sole suppliers. These sole suppliers, and any of our other suppliers or our third-party contract manufacturers, may be unwilling or unable to supply the necessary materials and components or manufacture and assemble our products reliably and at the levels we anticipate or that are required by the market. Our ability to supply our products commercially and to develop any future products depends, in part, on our ability to obtain these materials, components, and products in accordance with regulatory requirements and in sufficient quantities for commercialization and clinical testing. While our suppliers and contract manufacturers have generally met our demand for their products and services on a timely basis in the past, we cannot guarantee that they will in the future be able to meet our demand for their products, which could be adversely affected due to, for example, natural and man-made disasters, public health emergencies, pandemics such as COVID-19, other catastrophic events, the nature of our agreements with our contract manufacturers, our relative importance to such manufacturers as a customer or a contract manufacturer's decision to discontinue or reduce the level of business they conduct with us. If we are required to change contract manufacturers due to any change in or termination of our relationships with these third parties, or if our manufacturers are unable to obtain the materials they need to produce our products at consistent prices or at all, we may lose sales, experience manufacturing or other delays, incur increased costs or otherwise experience impairment to our customer relationships. We cannot guarantee that we will be able to establish alternative relationships on similar terms, without delay or at all.

While we believe replacement suppliers and manufacturers exist for all materials, components, and services necessary to manufacture our Inspire system, establishing additional or replacement suppliers for any of these materials, components or services, if required, could be time-consuming and expensive, may result in interruptions in our operations and product delivery, may affect the performance specifications of our Inspire system or could require that we modify its design. Even if we are able to find replacement suppliers or third-party contract manufacturers, we will be required to verify that the new supplier or third-party manufacturer maintains facilities, procedures, and operations that comply with our quality expectations and applicable regulatory requirements. Furthermore, our contract manufacturers could require us to move to another one of their production facilities or use alternative materials or components. Any of these events could require that we obtain a new regulatory authority approval before we implement the change, which could result in further delay and which may not be obtained at all. While we seek to maintain sufficient levels of inventory as discussed above, those inventories may not fully protect us from supply interruptions.

If our third-party suppliers fail to deliver the required commercial quantities of materials on a timely basis and at commercially reasonable prices, and we are unable to find one or more replacement suppliers capable of production at a substantially equivalent cost in substantially equivalent volumes and quality on a timely basis, the continued commercialization of our Inspire system, the supply of our products to customers, and the development of any future products will be delayed, limited or prevented, which could have material adverse effect on our business, financial condition, and results of operations.

***Performance issues, service interruptions or price increases by our shipping carriers could adversely affect our business and harm our reputation and ability to provide our services on a timely basis.***

Expedited, reliable shipping is essential to our operations. We rely heavily on providers of transport services for reliable and secure point-to-point transport of our Inspire system to our customers and for tracking of these shipments. Should a carrier encounter delivery performance issues such as loss, damage or destruction of any systems, such occurrences may damage our reputation and lead to decreased demand for our Inspire system and increased cost and expense to our business. Similarly, strikes, severe weather, natural disasters, public health crises or other service interruptions affecting delivery services we use would adversely affect our ability to process orders for our Inspire system on a timely basis.

***Consolidation in the healthcare industry or group purchasing organizations could lead to demands for price concessions, which may affect our ability to sell our products at prices necessary to support our current business strategies.***

Healthcare costs have risen significantly over the past decade, which has resulted in or led to numerous cost reform initiatives by legislators, regulators, and third-party payors. Cost reform has triggered a consolidation trend in the healthcare industry to aggregate purchasing power, which may create more requests for pricing concessions in the future. Additionally, group purchasing organizations, independent delivery networks, and large single accounts may continue to use their market power to consolidate purchasing decisions for hospitals and ASCs. We expect that market demand, government regulation, third-party coverage, and reimbursement policies and societal pressures will continue to change the healthcare industry worldwide, resulting in further business consolidations and alliances among our customers, which may exert further downward pressure on the prices of our products.

***We have limited experience marketing and selling our Inspire system, and if we are unable to expand, manage and maintain our direct sales and marketing organization we may not be able to generate revenue growth.***

We began selling our Inspire system in certain European markets in 2011, in the U.S. in 2014, and in Japan in 2021. As a result, we have limited experience marketing and selling our Inspire system. We currently sell our Inspire system through a direct sales force that targets ENT physicians and sleep centers in the U.S. and Europe, and also utilize various direct-to-consumer marketing initiatives, including paid online search, radio, television, social media, and online videos. In Japan, we sell our products through a distributor. As of December 31, 2021, our direct sales and marketing organization, including reimbursement personnel, consisted of 363 employees, having increased from 47 employees as of December 31, 2016. Our operating results are directly dependent upon the efforts of these employees. If our direct sales force fails to adequately promote, market and sell our Inspire system, our revenue may be adversely affected.

In order to generate future revenue growth, we plan to expand the size and geographic scope of our direct sales organization. This growth may require us to split or adjust existing sales territories, which may adversely affect our ability to retain customers in those territories. Additionally, our future success will depend largely on our ability to continue to hire, train, retain, and motivate skilled sales and reimbursement personnel with significant industry experience and technical knowledge of implantable devices and related products. Because the competition for their services is high, we cannot ensure that we will be able to hire and retain additional personnel on favorable or commercially reasonable terms, if at all. Failure to hire or retain qualified sales and reimbursement personnel would prevent us from expanding our business and generating revenue. If we are unable to expand our sales and marketing capabilities, we may not be able to effectively commercialize our Inspire system, which could have an adverse effect on our business, financial condition, and results of operations.

***To successfully market and sell our Inspire system in markets outside of the U.S., we must address many international business risks with which we have limited experience.***

Sales in markets outside of the U.S. accounted for approximately 5.3%, 8.0%, and 10.2% of our revenue for the years ended December 31, 2021, 2020, and 2019, respectively. Our strategy is to increase our international presence in Europe, including Germany and the Netherlands, as well as other international markets, such as Japan and Australia, which may increase our revenue from markets outside the U.S. International sales are subject to a number of risks, including:

- difficulties in staffing and managing our international operations;
- increased competition as a result of more products and procedures receiving regulatory approval or otherwise free to market in international markets;
- longer accounts receivable payment cycles and difficulties in collecting accounts receivable;
- reduced or varied protection for intellectual property rights in some countries;

- export restrictions, trade regulations, and foreign tax laws;
- fluctuations in currency exchange rates;
- foreign certification and regulatory clearance or approval requirements;
- difficulties in developing effective marketing campaigns in unfamiliar foreign countries;
- customs clearance and shipping delays;
- political, social, and economic instability abroad, terrorist attacks, and security concerns in general;
- preference for locally produced products;
- potentially adverse tax consequences, including the complexities of foreign value-added tax systems, tax inefficiencies related to our corporate structure, and restrictions on the repatriation of earnings;
- the burdens of complying with a wide variety of foreign laws and different legal standards; and
- increased financial accounting and reporting burdens and complexities.

If one or more of these risks are realized, our business, financial condition, and results of operations could be adversely affected.

***We primarily rely on our own direct sales force for our Inspire system, which may result in higher fixed costs than our competitors and may slow our ability to reduce costs in the face of a sudden decline in demand for our products.***

We primarily rely on our own direct sales force, which as of December 31, 2021, covered 157 territories in the U.S. and 12 in Europe, to market and sell our Inspire system. Some of our competitors rely predominantly on independent sales agents and third-party distributors. A direct sales force may subject us to higher fixed costs than those of companies that market competing products through independent third parties, due to the costs that we will bear associated with employee benefits, training, and managing sales personnel. As a result, we could be at a competitive disadvantage. Additionally, these fixed costs may slow our ability to reduce costs in the face of a sudden decline in demand for our Inspire system, which could have a material adverse effect on our business, financial condition, and results of operations.

***We may be unable to manage our growth effectively.***

Our past growth has provided, and our future growth may create, challenges for our organization. From December 31, 2016 to December 31, 2021, the number of our employees increased from 79 to 485. In the future, we expect to hire and train new personnel as we continue to grow and expand our operations. This growth may place significant strain on us. Successful growth is also dependent upon our ability to implement appropriate financial and management controls and systems and procedures. If we fail to manage these challenges effectively, there may be an adverse effect on our business, financial condition, and results of operations.

***Our ability to maintain our competitive position depends on our ability to attract and retain senior management and other highly qualified personnel.***

Our success depends in part on our continued ability to attract, retain, and motivate highly qualified management, clinical, and other personnel. We are highly dependent upon our management team, particularly our Chief Executive Officer and President and the rest of our senior management, and other key personnel. Although we have entered into employment letter agreements with all of our executive officers, each of them may terminate their employment with us at any time. The replacement of any of our key personnel likely would involve significant time and costs and may significantly delay or prevent the achievement of our business objectives and could therefore have an adverse effect on our business. Our future success also depends on our ability to continue to

attract and retain additional executive officers and other key employees who have the requisite background and skills relevant to our business and industry, a product like Inspire therapy and a company at our stage of growth.

***We face the risk of product liability claims that could be expensive, divert management's attention and harm our reputation and business. We may not be able to maintain adequate product liability insurance.***

Our business exposes us to the risk of product liability claims that are inherent in the testing, manufacturing and marketing of medical devices. This risk exists even if a device is cleared, approved, or certified for commercial sale by the FDA or foreign regulatory authorities or notified bodies and manufactured in facilities regulated by the FDA or an applicable foreign regulatory authority. Our Inspire system is designed to affect, and any future products will be designed to affect, important bodily functions and processes. Any side effects, manufacturing defects, misuse or abuse associated with our Inspire system could result in patient injury or death. The medical device industry has historically been subject to extensive litigation over product liability claims, and we cannot offer any assurance that we will not face product liability suits. We may be subject to product liability claims if our Inspire system causes, or is alleged to have caused, patient injury or death. In addition, an injury that is caused by the activities of our suppliers, such as those who provide us with components and raw materials, may be the basis for a claim against us. Product liability claims may be brought against us by patients, healthcare providers or others selling or otherwise coming into contact with our Inspire system, among others. If we cannot successfully defend ourselves against product liability claims, we will incur substantial liabilities and reputational harm. In addition, regardless of merit or eventual outcome, product liability claims may result in:

- costs of litigation;
- distraction of management's attention from our primary business;
- the inability to commercialize our Inspire system or new products;
- decreased demand for our Inspire system;
- damage to our business reputation;
- product recalls or withdrawals from the market;
- withdrawal of clinical trial participants;
- substantial monetary awards to patients or other claimants; or
- loss of sales.

While we may attempt to manage our product liability exposure by proactively recalling or withdrawing from the market any defective products, any recall or market withdrawal of our products may delay the supply of those products to our customers and may impact our reputation. We can provide no assurance that we will be successful in initiating appropriate market recall or market withdrawal efforts that may be required in the future or that these efforts will have the intended effect of preventing product malfunctions and the accompanying product liability that may result. Such recalls and withdrawals may also be used by our competitors to harm our reputation for safety or be perceived by patients as a safety risk when considering the use of our products, either of which could have a material adverse effect on our business, financial condition and results of operations.

Although we have product liability and clinical study liability insurance that we believe is appropriate, this insurance is subject to deductibles and coverage limitations. Our current product liability insurance may not continue to be available to us on acceptable terms, if at all, and, if available, coverage may not be adequate to protect us against any future product liability claims. If we are unable to obtain insurance at an acceptable cost or on acceptable terms or otherwise protect against potential product liability claims, we could be exposed to significant liabilities. A product liability claim, recall or other claim with respect to uninsured liabilities or for amounts in excess of insured liabilities could have a material adverse effect on our business, financial condition and results of operations.

***If the quality of our Inspire system does not meet the expectations of physicians or patients, then our brand and reputation or our business could be adversely affected.***

In the course of conducting our business, we must adequately address quality issues that may arise with our Inspire system, including defects in third-party components included in our Inspire system. Although we have established internal procedures designed to minimize risks that may arise from quality issues, there can be no assurance that we will be able to eliminate or mitigate occurrences of these issues and associated liabilities. In addition, even in the absence of quality issues, we may be subject to claims and liability if the performance of our Inspire system does not live up to the expectations of physicians or patients as a result of the patient's use of the product. For example, battery life will vary based on usage and therapy settings. Based on STAR trial therapy settings at the 12-month endpoint, the battery in our current generation neurostimulator is generally expected to last for approximately 11 years, but it may not last that long if a patient's use of the device or chosen level of stimulation is greater than expected. The minimum estimated longevity based on STAR trial results is seven years. If the quality of our Inspire system does not meet the expectations of physicians or patients, then our brand and reputation with those physicians or patients, or our business, financial condition and results of operations, could be adversely affected.

***If we choose to acquire new and complementary businesses, products or technologies, we may be unable to complete these acquisitions or to successfully integrate them in a cost-effective and non-disruptive manner.***

Our success depends, in part, on our ability to continually enhance and broaden our product offerings in response to changing customer demands, competitive pressures and advances in technologies. Accordingly, we may in the future pursue the acquisition of, or joint ventures relating to, complementary businesses, products or technologies instead of developing them ourselves. We do not know if we will be able to successfully complete any future acquisitions or joint ventures, or whether we will be able to successfully integrate any acquired business, product or technology or retain any key employees related thereto. Integrating any business, product or technology we acquire could be expensive and time-consuming, disrupt our ongoing business and distract our management. If we are unable to integrate any acquired businesses, products or technologies effectively, our business will be adversely affected. In addition, any amortization or charges resulting from the costs of acquisitions could increase our expenses.

***Unfavorable global economic conditions could adversely affect our business, financial condition or results of operations.***

Our results of operations could be adversely affected by general conditions in the global economy and in the global financial markets. The global financial crisis caused extreme volatility and disruptions in the capital and credit markets. Factors such as Brexit, the COVID-19 pandemic and the U.S. election cycles have contributed to this volatility. These global economic conditions could result in a variety of risks to our business, including weakened demand for our Inspire system, and adversely impact our ability to raise additional capital when needed on acceptable terms, if at all. A weak or declining economy could also strain our manufacturers or suppliers, possibly resulting in supply disruption, or cause our customers to delay making payments for our services. Any of the foregoing could harm our business and we cannot anticipate all of the ways in which the economic climate and financial market conditions could adversely affect our business.

***Failure of a key information technology system, process or site could have an adverse effect on our business.***

We rely extensively on information technology systems to conduct our business. These systems affect, among other things, ordering and managing materials from suppliers, shipping products to customers, processing transactions, summarizing and reporting results of operations, complying with regulatory, legal or tax requirements, data security and other processes necessary to manage our business. Our internal information technology systems and those of our third-party service providers, vendors, strategic partners and other contractors or consultants are vulnerable to damage or interruption from computer viruses, natural disasters, terrorism, war, telecommunication and electrical failures, hacking, cyberattacks, phishing attacks and other social engineering schemes, malicious code, employee theft or misuse, human error, fraud, denial or degradation of



service attacks, sophisticated nation-state and nation-state-supported actors or unauthorized access or use by persons inside our organization, or persons with access to systems inside our organization. The risk of a security breach or disruption, particularly through cyberattacks or cyber intrusion, including by computer hackers, foreign governments and cyber terrorists, has generally increased as the number, intensity and sophistication of attempted attacks and intrusions from around the world have increased and evolved. As a result of the COVID-19 pandemic, we and our third-party service providers and partners may also face increased cybersecurity risks due to our reliance on internet technology and the number of our employees who are working remotely, which may create additional opportunities for cybercriminals to exploit vulnerabilities. Furthermore, because the techniques used to obtain unauthorized access to, or to sabotage, systems change frequently and often are not recognized until launched against a target, we may be unable to anticipate these techniques or implement adequate preventative measures. We may experience security breaches that may remain undetected for an extended period. Our third-party service providers and partners are also subject to these heightened risks. If our systems are damaged or cease to function properly due to any number of causes, ranging from catastrophic events to power outages to security breaches, and our business continuity plans do not effectively compensate on a timely basis, we may experience interruptions in our operations, which could have an adverse effect on our business. Furthermore, any breach in our IT systems could lead to the unauthorized access, disclosure and use of non-public information, including information from our ADHERE patient registry or other patient information we create, receive, maintain or transmit, including with respect to our Inspire Cloud or the Inspire Sleep app, which is governed by HIPAA and other laws. Any such access, disclosure, or other loss of information could result in legal claims or proceedings, liability under laws that protect the privacy of personal information, and damage to our reputation.

In addition, we accept payments for our sales through credit and debit card transactions, which are handled through a third-party payment processor. As a result, we are subject to a number of risks related to credit and debit card payments. As a result of these transactions, we pay interchange and other fees, which may increase over time and could require us to either increase the prices we charge for our Inspire system or experience an increase in our costs and expenses. In addition, as part of the payment processing process, we transmit our customers' credit and debit card information to our third-party payment processor. We may in the future become subject to lawsuits or other proceedings for purportedly fraudulent transactions arising out of the actual or alleged theft of our customers' credit or debit card information if the security of our third-party credit card payment processor is breached. We and our third-party credit card payment processor are also subject to payment card association operating rules, certification requirements and rules governing electronic funds transfers, which could change or be reinterpreted to make it difficult or impossible for us to comply. If we or our third-party credit card payment processor fail to comply with these rules or requirements, we may be subject to fines and higher transaction fees and lose our ability to accept credit and debit card payments from our customers, and there may be an adverse effect on our business.

***If our facilities are damaged or become inoperable, we may be unable to continue to research, develop, and supply our Inspire system and, as a result, there could be an adverse effect on our business until we are able to secure a new facility and rebuild our inventory.***

We do not have redundant facilities. We perform substantially all of our research and development and back office activity at a single location in Golden Valley, Minnesota. The majority of our finished goods inventory is maintained at a third party location in Tennessee. Our facility, equipment and inventory would be costly to replace and could require substantial lead time to repair or replace. These facilities may be harmed or rendered inoperable by natural or man-made disasters, including, but not limited to, tornadoes, flooding, fire, power outages, and public health crises, which may render it difficult or impossible for us to perform our research, development and commercialization activities for some period of time. The inability to perform those activities, combined with the time it may take to rebuild our inventory of finished product, may result in the loss of customers or harm to our reputation. Although we possess insurance for damage to our property and the disruption of our business, this insurance may not be sufficient to cover all of our potential losses and this insurance may not continue to be available to us on acceptable terms, or at all.

***We rely on third-party distributors to effectively distribute our products in certain markets.***

We depend or expect to depend in the future on qualified distributors for the marketing and selling of our products in certain markets, namely in Japan, Singapore, and Hong Kong. If our distributors fail to effectively market and sell our Inspire system in full compliance with applicable laws, our operating results and business may suffer. Recruiting and retaining qualified third-party distributors and training them in our technology and product offering requires significant time and resources. To develop and expand our distribution, we may be required to scale and improve our processes and procedures that support our distributors. Further, if our relationship with a successful distributor terminates, we may be unable to replace that distributor without disruption to our business. If we fail to develop or maintain positive relationships with our distributors, including in new markets, fail to manage, train or incentivize these distributors effectively, or fail to provide distributors with competitive products on attractive terms, or if these distributors are not successful in their sales efforts, we may not achieve expected revenues or may have a reduction in revenue and our operating results, reputation and business would be harmed.

***We are subject to anti-bribery, anti-corruption, and anti-money laundering laws, including the U.S. Foreign Corrupt Practices Act, as well as export control laws, customs laws, sanctions laws and other laws governing our operations. If we fail to comply with these laws, we could be subject to civil or criminal penalties, other remedial measures and legal expenses, which could adversely affect our business, results of operations and financial condition.***

As we grow our international presence and global operations, we will have increasing obligations to comply with trade and economic sanctions and other restrictions imposed by the U.S., the European Union ("EU") and other governments and organizations. During the year ended December 31, 2021, approximately 5.3% of our total sales were made in EU member states and Japan. The U.S. Departments of Justice, Commerce, State and Treasury and other federal agencies and authorities have a broad range of civil and criminal penalties they may seek to impose against corporations and individuals for violations of economic sanctions laws, export control laws, the U.S. Foreign Corrupt Practices Act ("FCPA") and other federal statutes and regulations, including those established by the Office of Foreign Assets Control ("OFAC"). In addition, the U.K. Bribery Act of 2010 ("the Bribery Act") prohibits both domestic and international bribery, as well as bribery across both private and public sectors. An organization that "fails to prevent bribery" by anyone associated with the organization can be charged under the Bribery Act unless the organization can establish the defense of having implemented "adequate procedures" to prevent bribery. Under these laws and regulations, as well as other anti-corruption laws, anti-money laundering laws, export control laws, customs laws, sanctions laws and other laws governing our operations, various government agencies may require export licenses, may seek to impose modifications to business practices, including cessation of business activities in sanctioned countries or with sanctioned persons or entities and modifications to compliance programs, which may increase compliance costs, and may subject us to fines, penalties and other sanctions. A violation of these laws or regulations would negatively affect our business, financial condition and results of operations.

We have implemented policies and procedures designed to ensure compliance by us and our directors, officers, employees, representatives, consultants and agents with the FCPA, OFAC restrictions, the Bribery Act and other export control, anti-corruption, anti-money-laundering and anti-terrorism laws and regulations. We cannot ensure, however, that our policies and procedures are or will be sufficient or that directors, officers, employees, representatives, consultants and agents have not engaged and will not engage in conduct for which we may be held responsible, nor can we ensure that our business partners have not engaged and will not engage in conduct that could materially affect their ability to perform their contractual obligations to us or even result in our being held liable for such conduct. Violations of the FCPA, OFAC restrictions, the Bribery Act or other export control, anti-corruption, anti-money laundering and anti-terrorism laws or regulations may result in severe criminal or civil sanctions, and we may be subject to other liabilities, which could have a material adverse effect on our business, financial condition and results of operations.

***Our indebtedness may limit our flexibility in operating our business and adversely affect our financial health and competitive position.***

As of December 31, 2021, we had \$24.5 million of indebtedness outstanding under our credit facility with Oxford Finance LLC ("Oxford Finance"). In order to service this indebtedness and any additional indebtedness we may

incur in the future, we need to generate cash from our operating activities. Our ability to generate cash is subject, in part, to our ability to successfully execute our business strategy, as well as general economic, financial, competitive, regulatory and other factors beyond our control. We cannot ensure that our business will be able to generate sufficient cash flow from operations or that future borrowings or other financings will be available to us in an amount sufficient to enable us to service our indebtedness and fund our other liquidity needs. To the extent we are required to use cash from operations or the proceeds of any future financing to service our indebtedness instead of funding working capital, capital expenditures or other general corporate purposes, we will be less able to plan for, or react to, changes in our business, industry and in the economy generally. This will place us at a competitive disadvantage compared to our competitors that have less indebtedness.

In addition, the agreement governing the credit facility contains, and any agreements evidencing or governing other future indebtedness may contain, certain covenants that limit our ability to engage in certain transactions that may be in our long-term best interests. Subject to certain limited exceptions, these covenants limit our ability to, among other things:

- convey, sell, lease, transfer, assign, dispose of or otherwise make cash payments consisting of all or any part of our business or property;
- effect certain changes in our business, management, ownership or business locations;
- merge or consolidate with, or acquire all or substantially all of the capital stock or assets of, any other company;
- create, incur, assume or be liable for any additional indebtedness, or create, incur, allow or permit to exist any additional liens;
- pay cash dividends on, make any other distributions in respect of, or redeem, retire or repurchase, any shares of our capital stock;
- make certain investments; and
- enter into transactions with our affiliates.

While we have not previously breached and are not currently in breach of these or any of the other covenants contained in our credit agreement, there can be no guarantee that we will not breach these covenants in the future. Our ability to comply with these covenants may be affected by events and factors beyond our control. In the event that we breach one or more covenants, our lender may choose to declare an event of default and require that we immediately repay all amounts outstanding, terminate any commitment to extend further credit and foreclose on the collateral granted to it to collateralize such indebtedness. The occurrence of any of these events could have a material adverse effect on our business, financial condition and results of operations.

A portion of our indebtedness bears interest at variable interest rates, primarily based on the London Inter-bank Offered Rate ("LIBOR"), which may be subject to regulatory guidance and/or reform that could cause interest rates under our current or future debt agreements to perform differently than in the past or cause other unanticipated consequences. Some tenors of LIBOR were discontinued on December 31, 2021. Although we expect that the capital and debt markets will cease to use LIBOR as a benchmark in the near future and the administrator of LIBOR has announced its intention to extend the publication of most tenors of LIBOR for U.S. dollars through June 30, 2023, we cannot predict whether or when LIBOR will actually cease to be available, whether the Secured Overnight Funding Rate, or SOFR, will become the market benchmark in its place or what impact such a transition may have on our business, financial condition and results of operations.

***We bear the risk of warranty claims on our Inspire system.***

We bear the risk of warranty claims on our Inspire system. We may not be successful in claiming recovery under any warranty or indemnity provided to us by our suppliers or vendors in the event of a successful warranty claim against us by a customer or that any recovery from such vendor or supplier would be adequate. In addition,

warranty claims brought by our customers related to third-party components may arise after our ability to bring corresponding warranty claims against such suppliers expires, which could result in costs to us.

***We may need substantial additional funding beyond our existing cash resources and may be unable to raise capital when needed, which could force us to delay or reduce our commercialization efforts or product development programs.***

Our existing cash, cash equivalents, short-term investments and revenue will be sufficient to meet our capital requirements and fund our operations for at least 12 months. However, we have based these estimates on assumptions that may prove to be incorrect, and we could spend our available financial resources much faster than we currently expect. Any future funding requirements will depend on many factors, including:

- patient, physician and market acceptance of our Inspire therapy;
- the scope, rate of progress and cost of our current or future clinical studies;
- the cost of our research and development activities;
- the cost of filing and prosecuting patent applications and defending and enforcing our patent or other intellectual property rights;
- the cost of defending, in litigation or otherwise, any claims that we infringe third-party patents or other intellectual property rights;
- the cost and timing of additional regulatory clearances, approvals or certifications;
- the cost and timing of establishing additional sales and marketing capabilities;
- costs associated with any product recall that may occur;
- the effect of competing technological and market developments; and
- the extent to which we acquire or invest in products, technologies and businesses, although we currently have no commitments or agreements relating to any of these types of transactions.

Any additional equity or debt financing that we raise may contain terms that are not favorable to us or our stockholders. If we raise additional funds by selling additional shares of our common stock or other securities convertible into or exercisable or exchangeable for shares of our common stock, the issuance of such securities will result in dilution to our stockholders. The price per share at which we sell additional shares of our common stock, or securities convertible into or exercisable or exchangeable for shares of our common stock, in future transactions may be higher or lower than the price per share paid by investors in previous offerings of shares of our common stock. Furthermore, investors purchasing any securities we may issue in the future may have rights superior to the rights of a holder of our common stock.

In addition, any future debt financing into which we enter may impose upon us covenants that restrict our operations, including limitations on our ability to incur liens or additional debt, pay dividends, repurchase our common stock, make certain investments and engage in certain merger, consolidation or asset sale transactions. If we raise additional funds through collaboration and licensing arrangements with third-parties, it may be necessary to relinquish some rights to our technologies or our products, or grant licenses on terms that are not favorable to us.

Furthermore, we cannot be certain that additional funding will be available on acceptable terms, if at all. If we do not have, or are not able to obtain, sufficient funds, we may have to delay development or commercialization of our products or license to third-parties the rights to commercialize products or technologies that we would otherwise seek to commercialize. We also may have to reduce marketing, customer support or other resources

devoted to our products or cease operations. Any of these factors could harm our business, financial condition and results of operations.

***Our ability to use our net operating losses and research and development credit carryforwards to offset future taxable income may be subject to certain limitations.***

In general, under Sections 382 and 383 of the Internal Revenue Code of 1986, as amended ("the Code"), a corporation that undergoes an "ownership change," generally defined as a greater than 50% change by value in its equity ownership over a three-year period, is subject to limitations on its ability to utilize its pre-change net operating losses ("NOLs") and its research and development credit carryforwards to offset future taxable income. We have not performed a detailed analysis to determine whether an ownership change has occurred. As of December 31, 2021, our gross federal NOL carryforward was \$286.3 million. Our existing NOLs and research and development credit carryforwards may be subject to limitations arising from previous ownership changes, and if we undergo an ownership change, our ability to utilize NOLs and research and development credit carryforwards could be further limited by Sections 382 and 383 of the Code. In addition, our ability to deduct net interest expense may be limited if we have insufficient taxable income for the year during which the interest is incurred, and any carryovers of such disallowed interest would be subject to the limitation rules similar to those applicable to NOLs and other attributes. Future changes in our stock ownership, some of which might be beyond our control, could result in an ownership change under Section 382 of the Code. For these reasons, in the event we experience a change of control, we may not be able to utilize a material portion of the NOLs, research and development credit carryforwards or disallowed interest expense carryovers, even if we attain profitability.

***If we were deemed to be an investment company under the Investment Company Act of 1940, as amended (the "1940 Act"), applicable restrictions could make it impractical for us to continue our business as contemplated and could have a material adverse effect on our business, financial condition and results of operations.***

Under Sections 3(a)(1)(A) and (C) of the 1940 Act, a company generally will be deemed to be an "investment company" for purposes of the 1940 Act if (1) it is, or holds itself out as being, engaged primarily, or proposes to engage primarily, in the business of investing, reinvesting or trading in securities or (2) it engages, or proposes to engage, in the business of investing, reinvesting, owning, holding or trading in securities and it owns or proposes to acquire investment securities having a value exceeding 40% of the value of its total assets (exclusive of U.S. government securities and cash items) on an unconsolidated basis. We do not believe that we are an "investment company," as such term is defined in either of those sections of the 1940 Act.

We intend to conduct our operations so that we will not be deemed an investment company. However, if we were to be deemed an investment company, restrictions imposed by the 1940 Act, including limitations on our capital structure and our ability to transact with affiliates, could make it impractical for us to continue our business as contemplated and could have a material adverse effect on our business, financial condition and results of operations.

***The increasing focus on environmental sustainability and social initiatives could increase our costs, harm our reputation and adversely impact our financial results.***

There has been increasing public focus by investors, customers, environmental activists, the media, and governmental and nongovernmental organizations on a variety of environmental, social and other sustainability matters. We experience pressure to make commitments relating to sustainability matters that affect us, including the design and implementation of specific risk mitigation strategic initiatives relating to sustainability. If we are not effective in addressing environmental, social and other sustainability matters affecting our business, or setting and meeting relevant sustainability goals, our reputation and financial results may suffer. We may experience increased costs in order to execute upon our sustainability goals and measure achievement of those goals, which could have an adverse impact on our business and financial condition.

In addition, this emphasis on environmental, social and other sustainability matters has resulted and may result in the adoption of new laws and regulations, including new reporting requirements. If we fail to comply with new laws, regulations or reporting requirements, our reputation and business could be adversely impacted.

## **Risks Related to Government Regulation**

***Our products and operations are subject to extensive government regulation and oversight both in the U.S. and abroad, and our failure to comply with applicable requirements could harm our business.***

We and our products are subject to extensive regulation in the U.S. and elsewhere, including by the FDA and its foreign counterparts. The FDA and foreign regulatory agencies regulate, among other things, with respect to medical devices: design, development and manufacturing; testing, labeling, content and language of instructions for use and storage; clinical trials; product safety; establishment registration and device listing; marketing, sales and distribution; pre-market clearance, approval, and certification; record keeping procedures; advertising and promotion; recalls and field safety corrective actions; post-market surveillance, including reporting of deaths or serious injuries and malfunctions that, if they were to recur, could lead to death or serious injury; post-market approval studies; and product import and export.

The regulations to which we are subject are complex and have tended to become more stringent over time. Regulatory changes could result in restrictions on our ability to carry on or expand our operations, higher than anticipated costs or lower than anticipated sales. The FDA and foreign regulatory authorities enforce these regulatory requirements through periodic unannounced inspections. Failure to comply with applicable regulations could jeopardize our ability to sell our products and result in enforcement actions such as: warning letters; fines; injunctions; civil penalties; termination of distribution; recalls or seizures of products; delays in the introduction of products into the market; total or partial suspension of production; refusal to grant future clearances, approval, or certifications; withdrawals or suspensions of current approvals, resulting in prohibitions on sales of our products; and in the most serious cases, criminal penalties.

***We may not receive the necessary approvals or certifications for our future products or expanded indications, and failure to timely obtain necessary approvals or certifications for our future products or expanded indications would adversely affect our ability to grow our business.***

An element of our strategy is to continue to upgrade our products, add new features and expand the indications and uses for our current products. In the U.S., before we can market a new medical device, or a new use of, or claim for, or significant modification to, an existing product, we must first receive PMA from the FDA. In the process of obtaining PMA, which was required for our Inspire system, the FDA must determine that a proposed device is safe and effective for its intended use based, in part, on extensive data, including, but not limited to, technical, pre-clinical, clinical trial, manufacturing and labeling data. The PMA process is typically required for devices that are deemed to pose the greatest risk, such as life-sustaining, life-supporting or implantable devices.

Modifications to products that are approved through a PMA application generally require FDA approval. PMA can be expensive, lengthy and uncertain. The process of obtaining a PMA is costly and more uncertain than the 510(k) clearance process used for lower risk devices. PMAs generally take from one to three years, or even longer, from the time the application is filed with the FDA. In addition, a PMA often requires the performance of one or more clinical trials. Despite the time, effort and cost, a device may not be approved by the FDA. Any delay or failure to obtain necessary regulatory approvals could harm our business. Furthermore, even if we are granted regulatory approval, it may include significant limitations on the indicated uses for the device, which may limit the market for the device.

In the U.S., we have obtained approval of our Inspire system through the PMA pathway. Any modification to the Inspire system that has not been previously approved may require us to submit a new PMA or PMA supplement and obtain FDA approval prior to implementing the change. If the FDA requires us to go through a lengthier, more rigorous examination for future products or modifications to existing products than we had expected, product introductions or modifications could be delayed or canceled, which could adversely affect our ability to grow our business.

The FDA and other regulatory authorities or notified bodies outside the U.S. can delay, limit or deny approval or certification of a device for many reasons, including:

- our inability to demonstrate to the satisfaction of the FDA or the applicable foreign regulatory entity or notified body that our products are safe or effective for their intended uses;
- the disagreement of the FDA or the applicable foreign regulatory authority or notified body with the design or implementation of our clinical trials or the interpretation of data from pre-clinical studies or clinical trials;
- serious and unexpected adverse device effects experienced by participants in our clinical trials;
- the data from our pre-clinical studies and clinical trials may be insufficient to support approval or certification, where required;
- our inability to demonstrate that the clinical and other benefits of the device outweigh the risks;
- the manufacturing process or facilities we use may not meet applicable requirements; and
- the potential for approval policies or regulations of the FDA or applicable foreign regulatory authorities to change significantly in a manner rendering our clinical data or regulatory filings insufficient for clearance, approval or certification.

In addition, the FDA may change its approval policies, adopt additional regulations or revise existing regulations, or take other actions, which may prevent or delay approval of our future products under development or impact our ability to modify our currently cleared products on a timely basis. Such policy or regulatory changes could impose additional requirements upon us that could delay our ability to obtain new approvals, increase the costs of compliance or restrict our ability to maintain our current approval.

The EU landscape concerning medical devices recently evolved. On May 25, 2017, the EU Medical Devices Regulation entered into force, which repeals and replaces the Council Directive 93/42/EEC (the “EU Medical Devices Directive”) and Directive 90/385/EEC (“AIMDD”). Unlike directives, which must be implemented into the national laws of the EU member states, regulations are directly applicable (i.e., without the need for adoption of EU member state laws implementing them) in all EU member states and are intended to eliminate current differences in the regulation of medical devices among EU member states. Devices lawfully placed on the market pursuant to the EU Medical Devices Directive or the AIMDD prior to May 26, 2021 may generally continue to be made available on the market or put into service until May 26, 2025, provided that the requirements of the transitional provisions are fulfilled. In particular, the certificate in question must still be valid. However, even in this case, manufacturers must comply with a number of new or reinforced requirements set forth in the EU Medical Devices Regulation with regard to registration of economic operators and of devices, post-market surveillance, market surveillance and vigilance requirements.

Subject to the transitional provisions, in order to sell our products in EU member states, our products must comply with the general safety and performance requirements of the EU Medical Devices Regulation, which repeals and replaces EU Medical Devices Directive and the AIMDD. Compliance with these requirements is a prerequisite to be able to affix the European Conformity (“CE”) mark to our products, without which they cannot be sold or marketed in the EU. See — Government Regulation. To demonstrate compliance with the general safety and performance requirements, we must undergo a conformity assessment procedure, which varies according to the type of medical device and its (risk) classification. Except for low risk medical devices (Class I), where the manufacturer can self-assess the conformity of its products with the general safety and performance requirements (except for any parts which relate to sterility, metrology or reuse aspects), a conformity assessment procedure requires the intervention of a notified body. The notified body would typically audit and examine the technical file and the quality system for the manufacture, design and final inspection of our devices. If satisfied that the relevant product conforms to the relevant general safety and performance requirements, the notified body issues a certificate of conformity, which the manufacturer uses as a basis for its own declaration of conformity. The manufacturer may then apply the CE mark to the device, which allows the device to be placed on the market throughout the EU. If we fail to comply with applicable laws and regulations, we would be unable to affix the CE mark to our products, which would prevent us from selling them within the EU.

We must inform the notified body that carried out the conformity assessment of the medical devices that we market or sell in the EU and the EEA of any planned substantial changes to our quality system or substantial changes to our medical devices that could affect compliance with the general safety and performance requirements laid down in Annex I to the EU Medical Devices Regulation or cause a substantial change to the intended use for which the device has been CE marked. The notified body will then assess the planned changes and verify whether they affect the products' ongoing conformity with the EU Medical Devices Regulation. If the assessment is favorable, the notified body will issue a new certificate of conformity or an addendum to the existing certificate attesting compliance with the general safety and performance requirements and quality system requirements laid down in the Annexes to the EU Medical Devices Regulation.

***Modifications to our products may require us to obtain new PMAs or approvals of a PMA supplement, and if we market modified products without obtaining necessary approvals, we may be required to cease marketing or recall the modified products until required approvals are obtained.***

Certain modifications to a PMA-approved device may require approval of a new PMA or a PMA supplement, or alternatively a notification or other submission to the FDA. The FDA may not agree with our decisions regarding whether a new PMA or PMA supplement is necessary. We may make modifications to our approved devices in the future that we believe do not require approval of a new PMA or PMA supplement. If the FDA disagrees with our determination and requires us to submit a new PMA or PMA supplement for modifications to our previously approved products, we may be required to cease marketing or to recall the modified product until we obtain approval, and we may be subject to significant regulatory fines or penalties. In addition, the FDA may not approve our products for the indications that are necessary or desirable for successful commercialization or could require clinical trials to support any modifications. Similar requirements may apply in foreign jurisdictions where we market our products. Any delay or failure in obtaining required approvals or certifications would adversely affect our ability to introduce new or enhanced products in a timely manner, which in turn would harm our future growth.

***Failure to comply with post-marketing regulatory requirements could subject us to enforcement actions, including substantial penalties, and might require us to recall or withdraw a product from the market.***

Even though we have obtained approval for the Inspire system, we are subject to ongoing and pervasive regulatory requirements governing, among other things, the manufacture, marketing, advertising, medical device reporting, sale, promotion, registration, and listing of devices. For example, we must submit periodic reports to the FDA as a condition of PMA. These reports include safety and effectiveness information about the device after its approval. Failure to submit such reports, or failure to submit the reports in a timely manner, could result in enforcement action by the FDA. Following its review of the periodic reports, the FDA might ask for additional information or initiate further investigation. Similar requirements may apply in foreign jurisdictions where we market our products.

In addition, the PMA for our Inspire system was subject to several conditions of approval, including a post-market long-term study. Though we believe we have complied with these conditions to date, any failure to comply with the conditions of approval could result in the withdrawal of PMA and the inability to continue to market the device. Failure to conduct the required studies in accordance with institutional review board ("IRB") and informed consent requirements, or adverse findings in these studies, could also be grounds for withdrawal of approval of the PMA.

The regulations to which we are subject are complex and have become more stringent over time. Regulatory changes could result in restrictions on our ability to continue or expand our operations, higher than anticipated costs, or lower than anticipated sales. Even after we have obtained the proper regulatory approval or certification to market a device, we have ongoing responsibilities under FDA regulations and applicable foreign laws and regulations. The FDA, state and foreign regulatory authorities have broad enforcement powers. Our failure to comply with applicable regulatory requirements could result in enforcement action by the FDA, state or foreign regulatory authorities, which may include any of the following sanctions:

- untitled letters or warning letters;
- fines, injunctions, consent decrees and civil penalties;



- recalls, termination of distribution, administrative detention, or seizure of our products;
- customer notifications or repair, replacement or refunds;
- operating restrictions or partial suspension or total shutdown of production;
- delays in or refusal to grant our requests for future PMAs or foreign regulatory approvals or certifications of new products, new intended uses, or modifications to existing products;
- withdrawals or suspensions of our current PMA or foreign regulatory approvals or certifications, resulting in prohibitions on sales of our products;
- FDA refusal to issue certificates to foreign governments needed to export products for sale in other countries; and
- criminal prosecution.

Any of these sanctions could result in higher than anticipated costs or lower than anticipated sales and have a material adverse effect on our reputation, business, financial condition and results of operations.

***Our products must be manufactured in accordance with foreign, federal and state regulations, and we or any of our suppliers or third-party manufacturers could be forced to recall our installed systems or terminate production if we fail to comply with these regulations.***

The methods used in, and the facilities used for, the manufacture of our products must comply with the FDA's Quality System Regulation ("QSR") which is a complex regulatory scheme that covers the procedures and documentation of the design, testing, production, process controls, quality assurance, labeling, packaging, handling, storage, distribution, installation, servicing and shipping of medical devices. Furthermore, we are required to verify that our suppliers maintain facilities, procedures and operations that comply with our quality standards and applicable regulatory requirements. The FDA enforces the QSR through periodic announced or unannounced inspections of medical device manufacturing facilities, which may include the facilities of subcontractors. Our products are also subject to similar state regulations and various laws and regulations of foreign countries governing manufacturing.

Our third-party manufacturers may not take the necessary steps to comply with applicable regulations, which could cause delays in the delivery of our products. In addition, failure to comply with applicable FDA foreign requirements or later discovery of previously unknown problems with our products or manufacturing processes could result in, among other things: warning letters or untitled letters; fines, injunctions or civil penalties; suspension or withdrawal of approvals or certifications; seizures or recalls of our products; total or partial suspension of production or distribution; administrative or judicially imposed sanctions; the FDA's or applicable foreign regulatory authority's or notified body's refusal to grant pending or future clearances, approvals or certifications for our products; clinical holds; refusal to permit the import or export of our products; and criminal prosecution of us or our employees.

Any of these actions could significantly and negatively affect supply of our products. If any of these events occurs, our reputation could be harmed, we could be exposed to product liability claims and we could lose customers and experience reduced sales and increased costs.

***If treatment guidelines for OSA change or the standard of care evolves, we may need to redesign and seek new marketing authorization from the FDA for one or more of our products.***

If treatment guidelines for OSA changes or the standard of care for this condition evolves, we may need to redesign the applicable product and seek new approvals from the FDA. Our PMAs from the FDA are based on current treatment guidelines. If treatment guidelines change so that different treatments become desirable, the clinical utility of one or more of our products could be diminished and our business could be adversely affected.

***The misuse or off-label use of our Inspire system may harm our reputation in the marketplace, result in injuries that lead to product liability suits or result in costly investigations, fines or sanctions by regulatory bodies if we are deemed to have engaged in the promotion of these uses, any of which could be costly to our business.***

Our Inspire system has been approved by the FDA for specific indications. We train our marketing personnel and direct sales force to not promote our Inspire system for uses outside of the FDA-approved indications for use, known as “off-label uses.” We cannot, however, prevent a physician from using our Inspire system off-label, when in the physician’s independent professional medical judgment he or she deems it appropriate. There may be increased risk of injury to patients if physicians attempt to use our Inspire system off-label. Furthermore, the use of our Inspire system for indications other than those approved by the FDA, approved by any foreign regulatory authority or certified by a notified body, may not effectively treat such conditions, which could harm our reputation in the marketplace among physicians and patients.

If the FDA or any foreign regulatory authority determines that our promotional materials or training constitute promotion of an off-label use, it could request that we modify our training or promotional materials or subject us to regulatory or enforcement actions, including the issuance or imposition of an untitled letter, which is used for violators that do not necessitate a warning letter, injunction, seizure, civil fine or criminal penalties. It is also possible that other federal, state or foreign enforcement authorities might take action under other regulatory authority, such as false claims laws, if they consider our business activities to constitute promotion of an off-label use, which could result in significant penalties, including, but not limited to, criminal, civil and administrative penalties, damages, fines, disgorgement, exclusion from participation in government healthcare programs and the curtailment of our operations.

In addition, physicians may misuse our Inspire system or use improper techniques if they are not adequately trained, potentially leading to injury and an increased risk of product liability. If our Inspire system is misused or used with improper technique, we may become subject to costly litigation by our customers or their patients. Similarly, in an effort to decrease costs, physicians may also reuse our Inspire system despite it being intended for a single use or may purchase reprocessed Inspire systems from third-party reproducers in lieu of purchasing a new Inspire system from us, which could result in product failure and liability. As described above, product liability claims could divert management’s attention from our core business, be expensive to defend and result in sizable damage awards against us that may not be covered by insurance.

***Our products may cause or contribute to adverse medical events or be subject to failures or malfunctions that we are required to report to the FDA and foreign regulatory authorities, and if we fail to do so, we would be subject to sanctions that could harm our reputation, business, financial condition and results of operations. The discovery of serious safety issues with our products, or a recall of our products either voluntarily or at the direction of the FDA or another governmental authority, could have a negative impact on us.***

We are subject to the FDA’s medical device reporting regulations and similar foreign regulations, which require us to report to the FDA and foreign regulatory authorities when we receive or become aware of information that reasonably suggests that one or more of our products may have caused or contributed to a death or serious injury or malfunctioned in a way that, if the malfunction were to recur, it could cause or contribute to a death or serious injury. The timing of our obligation to report is triggered by the date we become aware of the adverse event as well as the nature of the event. We may fail to report adverse events of which we become aware within the prescribed timeframe. We may also fail to recognize that we have become aware of a reportable adverse event, especially if it is not reported to us as an adverse event or if it is an adverse event that is unexpected or removed in time from the use of the product. If we fail to comply with our reporting obligations, the FDA and foreign regulatory authorities could take action, including warning letters, untitled letters, administrative actions, criminal prosecution, imposition of civil monetary penalties, revocation of our device approval or certification, seizure of our products or delay in clearance, approval or certification of future products.

The FDA and foreign regulatory bodies have the authority to require the recall of commercialized products in the event of material deficiencies or defects in design or manufacture of a product or in the event that a product poses an unacceptable risk to health. The FDA’s and foreign regulatory bodies’ authority to require a recall must be

based on a finding that there is reasonable probability that the device could cause serious injury or death. We may also choose to voluntarily recall a product if any material deficiency is found. A government-mandated or voluntary recall by us could occur as a result of an unacceptable risk to health, component failures, malfunctions, manufacturing defects, labeling or design deficiencies, packaging defects or other deficiencies or failures to comply with applicable regulations. Product defects or other errors may occur in the future.

Depending on the corrective action we take to redress a product's deficiencies or defects, the FDA or foreign regulatory authorities may require, or we may decide, that we will need to obtain new approvals for the device before we may market or distribute the corrected device. Seeking such approvals may delay our ability to replace the recalled devices in a timely manner. Moreover, if we do not adequately address problems associated with our devices, we may face additional regulatory enforcement action, including FDA or foreign regulatory authority warning letters, product seizure, injunctions, administrative penalties or civil or criminal fines.

Companies are required to maintain certain records of recalls and corrections, even if they are not reportable to the FDA or foreign regulatory authorities. We may initiate voluntary withdrawals or corrections for our products in the future that we determine do not require notification to the FDA or foreign regulatory authorities. If the FDA or a foreign regulatory authority disagrees with our determinations, it could require us to report those actions as recalls and we may be subject to enforcement action. A future recall announcement could harm our reputation with customers, potentially lead to product liability claims against us and negatively affect our sales. Any corrective action, whether voluntary or involuntary, as well as defending ourselves in a lawsuit, will require the dedication of our time and capital, distract management from operating our business and may harm our reputation and financial results.

***If we do not obtain and maintain international regulatory registrations, approvals or certifications for our products, we will be unable to market and sell our products outside of the U.S.***

Sales of our products outside of the U.S. are subject to foreign regulatory requirements that vary widely from country to country. In addition, the FDA regulates exports of medical devices from the U.S. While the regulations of some countries may not impose barriers to marketing and selling our products or only require notification, others require that we obtain the approval of or certification by a specified body (e.g., notified bodies in Europe). Complying with foreign regulatory requirements, including obtaining registrations, approvals or certifications, can be expensive and time-consuming, and we may not receive regulatory approvals or certifications in each country in which we plan to market our products or we may be unable to do so on a timely basis. The time required to obtain registrations, approvals or certifications, if required by other countries, may be longer than that required for FDA approval, and requirements for such registrations, clearances, approvals or certifications may significantly differ from FDA requirements. If we modify our products, we may need to apply for additional regulatory approvals or certifications before we are permitted to sell the modified product. In addition, we may not continue to meet the quality and safety standards required to maintain the authorizations or certifications that we have received. If we are unable to maintain our authorizations or certifications in a particular country, we will no longer be able to sell the applicable product in that country.

Regulatory approval by the FDA does not ensure registration, clearance, approval or certification by regulatory authorities or notified bodies in other countries, and registration, clearance, approval or certification by one or more foreign regulatory authorities or notified bodies does not ensure registration, clearance, approval or certification by regulatory authorities or notified bodies in other foreign countries or by the FDA. However, a failure or delay in obtaining registration, regulatory clearance, approval or certification in one country may have a negative effect on the regulatory process in others.

***Legislative or regulatory reforms in the U.S. or the EU may make it more difficult and costly for us to obtain regulatory clearances, approvals or certification for our products or to manufacture, market or distribute our products after clearance, approval or certification is obtained.***

From time to time, legislation is drafted and introduced in Congress that could significantly change the statutory provisions governing the regulation of medical devices. 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 statutes, regulations or revisions or reinterpretations of existing regulations may impose additional costs or

lengthen review times of any future products or make it more difficult to obtain approval for, manufacture, market or distribute our products. We cannot determine what effect changes in regulations, statutes, legal interpretation or policies, when and if promulgated, enacted or adopted may have on our business in the future. Such changes could, among other things, require: additional testing prior to obtaining clearance or approval; changes to manufacturing methods; recall, replacement or discontinuance of our products; or additional record keeping.

Similarly, the EU landscape concerning medical devices recently evolved. On May 25, 2017, the EU Medical Devices Regulation entered into force, which repeals and replaces the EU Medical Devices Directive and the AIMDD. See Part I, Item I, "Business – Government Regulation" for additional information on these reforms. These modifications are likely to have an effect on the way we conduct our business in the EEA. For example, as a result of the transition towards the new regime, notified body review times have lengthened, and product introductions or modifications could be delayed or canceled, which could adversely affect our ability to grow our business.

***We are subject to certain federal, state and foreign fraud and abuse laws, health information privacy and security laws and transparency laws, which, if violated, could subject us to substantial penalties. Additionally, any challenge to or investigation into our practices under these laws could cause adverse publicity and be costly to respond to, and thus could harm our business.***

There are numerous U.S. federal and state, as well as foreign, laws pertaining to healthcare fraud and abuse, including anti-kickback, false claims and physician transparency laws. Our business practices and relationships with providers are subject to scrutiny under these laws. The healthcare laws and regulations that may affect our ability to operate include, but are not limited to:

- the federal Anti-Kickback Statute, which prohibits, among other things, persons and entities from knowingly and willfully soliciting, offering, receiving or providing remuneration, directly or indirectly, in cash or in kind, to induce either the referral of an individual or furnishing or arranging for a good or service, for which payment may be made, in whole or in part, under federal healthcare programs, such as Medicare and Medicaid. The U.S. government has interpreted this law broadly to apply to the marketing and sales activities of manufacturers. In addition, a person or entity does not need to have actual knowledge of the statute or specific intent to violate it to have committed a violation;
- the federal civil and criminal false claims laws and civil monetary penalties laws, including the federal civil False Claims Act, which prohibit, among other things, individuals or entities from knowingly presenting, or causing to be presented, claims for payment from Medicare, Medicaid or other federal healthcare programs that are false or fraudulent. Moreover, 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 civil False Claims Act. Private individuals can bring False Claims Act "qui tam" actions, on behalf of the government and such individuals, commonly known as "whistleblowers," may share in amounts paid by the entity to the government in fines or settlement;
- the federal Civil Monetary Penalties Law, which prohibits, among other things, offering or transferring remuneration to a federal healthcare beneficiary that a person knows or should know is likely to influence the beneficiary's decision to order or receive items or services reimbursable by the government from a particular provider or supplier;
- HIPAA, which created additional federal criminal statutes that prohibit, among other things, executing a scheme to defraud any healthcare benefit program and making false statements relating to healthcare matters. Similar to the federal Anti-Kickback Statute, a person or entity does not need to have actual knowledge of the statute or specific intent to violate it to have committed a violation;
- the federal Physician Payments Sunshine Act, which requires certain applicable manufacturers of drugs, devices, biologics and medical supplies for which payment is available under Medicare, Medicaid or the Children's Health Insurance Program ("CHIP") to report annually to the DHHS Centers for Medicare and Medicaid Services ("CMS") information related to payments and other transfers of value to physicians (as defined by statute), certain non-physician practitioners such as physician assistants and nurse

practitioners, and teaching hospitals, and applicable manufacturers and group purchasing organizations, to report annually ownership and investment interests held by physicians and their immediate family members; and

- analogous state and foreign law equivalents of each of the above federal laws, such as anti-kickback and false claims laws which may apply to items or services reimbursed by any third-party payor, including commercial insurers or patients; state laws that require device companies to comply with the industry's voluntary compliance guidelines and the applicable compliance guidance promulgated by the federal government or otherwise restrict payments that may be made to healthcare providers and other potential referral sources; state laws that require device manufacturers to report information related to payments and other transfers of value to physicians and other healthcare providers or marketing expenditures; consumer protection and unfair competition laws, which broadly regulate marketplace activities and activities that potentially harm customers; and state laws related to insurance fraud in the case of claims involving private insurers.

These laws and regulations, among other things, constrain our business, marketing and other promotional activities by limiting the kinds of financial arrangements, including sales programs, we may have with hospitals, physicians or other potential purchasers of our products. Similar laws may exist in other jurisdictions where we operate, such as in the EU. Due to the breadth of these laws, the narrowness of statutory exceptions and regulatory safe harbors available, and the range of interpretations to which they are subject, it is possible that some of our current or future practices might be challenged under one or more of these laws.

To enforce compliance with the healthcare regulatory laws, certain enforcement bodies have recently increased their scrutiny of interactions between healthcare companies and healthcare providers, which has led to a number of investigations, prosecutions, convictions and settlements in the healthcare industry. Responding to investigations can be time-and resource-consuming and can divert management's attention from the business. Additionally, as a result of these investigations, healthcare providers and entities may have to agree to additional compliance and reporting requirements as part of a consent decree or corporate integrity agreement. Any such investigation or settlement could increase our costs or otherwise have an adverse effect on our business. Even an unsuccessful challenge or investigation into our practices could cause adverse publicity, and be costly to respond to. If our operations are found to be in violation of any of the healthcare laws or regulations described above or any other healthcare regulations that apply to us, we may be subject to penalties, including administrative, civil and criminal penalties, damages, fines, exclusion from participation in government healthcare programs, such as Medicare and Medicaid, imprisonment, contractual damages, reputational harm, disgorgement and the curtailment or restructuring of our operations.

***We are subject to U.S. federal, state, and foreign laws and regulations imposing obligations on how we collect, store and process personal information. Our actual or perceived failure to comply with such obligations could harm our business. Ensuring compliance with such laws could also impair our efforts to maintain and expand our customer base, and thereby decrease our revenue.***

In the conduct of our business, we process health-related and other personal information. The U.S. federal government and various states have adopted or proposed laws, regulations, guidelines and rules for the collection, distribution, use and storage of personal information of individuals. For example, HIPAA, as amended by the Health Information Technology for Economic and Clinical Health Act of 2009, and regulations implemented thereunder (collectively "HIPAA"), imposes privacy, security and breach notification obligations on certain healthcare providers, health plans, and healthcare clearinghouses, known as covered entities, as well as their business associates that perform certain services that involve creating, receiving, maintaining or transmitting individually identifiable health information for or on behalf of such covered entities, and their covered subcontractors. HIPAA requires covered entities and business associates to develop and maintain policies with respect to the protection of, use and disclosure of PHI, including the adoption of administrative, physical and technical safeguards to protect such information, and certain notification requirements in the event of a breach of unsecured PHI.

Under HIPAA, covered entities must report breaches of unsecured PHI to affected individuals without unreasonable delay, not to exceed 60 days following discovery of the breach by a covered entity or its agents.

Notification also must be made to the U.S. Department of Health and Human Services Office for Civil Rights, or OCR, and, in certain circumstances involving large breaches, to the media. Business associates must report breaches of unsecured PHI to covered entities within 60 days of discovery of the breach by the business associate or its agents. A non-permitted use or disclosure of PHI is presumed to be a breach under HIPAA unless the covered entity or business associate establishes that there is a low probability the information has been compromised consistent with requirements enumerated in HIPAA.

Entities that are found to be in violation of HIPAA as the result of a breach of unsecured PHI, a complaint about privacy practices or an audit by the U.S. Department of Health and Human Services, or HHS, may be subject to significant civil, criminal and administrative fines and penalties and/or additional reporting and oversight obligations if required to enter into a resolution agreement and corrective action plan with HHS to settle allegations of HIPAA non-compliance. HIPAA also authorizes state Attorneys General to file suit on behalf of their residents. Courts may award damages, costs and attorneys' fees related to violations of HIPAA in such cases. While HIPAA does not create a private right of action allowing individuals to sue us in civil court for violations of HIPAA, its standards have been used as the basis for duty of care in state civil suits such as those for negligence or recklessness in the misuse or breach of PHI.

Further, the Federal Trade Commission (the "FTC") and many state Attorneys General continue to enforce federal and state consumer protection laws against companies for online collection, use, dissemination and security practices that appear to be unfair or deceptive. For example, according to the FTC, failing to take appropriate steps to keep consumers' personal information secure can constitute unfair acts or practices in or affecting commerce in violation of Section 5(a) of the Federal Trade Commission Act. The FTC expects a company's data security measures to be reasonable and appropriate in light of the sensitivity and volume of consumer information it holds, the size and complexity of its business, and the cost of available tools to improve security and reduce vulnerabilities.

We may also be subject to U.S. federal rules, regulations and guidance concerning data security for medical devices, including guidance from the FDA. State privacy and security laws vary from state to state and, in some cases, can impose more restrictive requirements than U.S. federal law. Where state laws are more protective, we must comply with the stricter provisions. In addition to fines and penalties that may be imposed for failure to comply with state law, some states also provide for private rights of action to individuals for misuse of personal information. For example, the California Consumer Privacy Act of 2018 (the "CCPA") went into effect on January 1, 2020. The CCPA creates individual privacy rights for California consumers and increases the privacy and security obligations of entities handling certain personal information. The CCPA provides for civil penalties for violations, as well as a private right of action for data breaches that is expected to increase data breach litigation. Further, the California Privacy Rights Act (the "CPRA") recently passed in California. The CPRA will impose additional data protection obligations on covered businesses, including additional consumer rights processes, limitations on data uses, new audit requirements for higher risk data, and opt outs for certain uses of sensitive data. It will also create a new California data protection agency authorized to issue substantive regulations and could result in increased privacy and information security enforcement. The majority of the provisions will go into effect on January 1, 2023, and additional compliance investment and potential business process changes may be required. Similar laws have passed in Virginia and Colorado, and have been proposed in other states and at the federal level, reflecting a trend toward more stringent privacy legislation in the United States. The enactment of such laws could have potentially conflicting requirements that would make compliance challenging.

In Europe, we are subject to the requirements of the GDPR (and national laws implementing the GDPR) because we are "established" in certain EU countries and we are processing personal data of individuals located in the EU and EEA in the context of these establishments, as well as offering of goods to, and/or monitoring the behavior of, individuals in the EU and EEA in connection with our clinical investigations. The GDPR, which went into effect in May 2018, imposes strict requirements for processing the personal data of individuals within the EEA. If we do not comply with our obligations under the GDPR, we could be exposed to significant fines the greater of EUR 20 million or 4% of total global annual turnover for certain breaches. In addition to the foregoing, a breach of the GDPR could result in regulatory investigations, reputational damage, orders to cease/ change our use of data, enforcement notices, as well potential civil claims including class action type litigation where individuals suffer harm.

Among other requirements, the GDPR regulates transfers of personal data subject to the GDPR to third countries that have not been found to provide adequate protection to such personal data, including the United States, and the efficacy and longevity of current transfer mechanisms between the EU and the United States remains uncertain. For example, in July 2020, the Court of Justice of the EU ("CJEU") limited how organizations could lawfully transfer personal data from the EU/EEA to the United States by invalidating the Privacy Shield for purposes of international transfers and imposing further restrictions on the use of standard contractual clauses, or SCCs. The European Commission issued revised SCCs on June 4, 2021 to account for the decision of the CJEU and recommendations made by the European Data Protection Board. The revised SCCs must be used for relevant new data transfers from September 27, 2021; existing standard contractual clauses arrangements must be migrated to the revised clauses by December 27, 2022. The new SCCs apply only to the transfer of personal data outside of the EEA and not the UK; the UK's Information Commissioner's Office launched a public consultation on its draft revised data transfers mechanisms in August 2021. There is some uncertainty around whether the revised clauses can be used for all types of data transfers, particularly whether they can be relied on for data transfers to non-EEA entities subject to the GDPR. As supervisory authorities issue further guidance on personal data export mechanisms, including circumstances where the SCCs cannot be used, and/or start taking enforcement action, we could suffer additional costs, complaints and/or regulatory investigations or fines, and/or if we are otherwise unable to transfer personal data between and among countries and regions in which we operate, it could affect the manner in which we provide our services, the geographical location or segregation of our relevant systems and operations, and could adversely affect our financial results

Further, from January 1, 2021, we have to comply with both the GDPR and the GDPR as incorporated into United Kingdom national law, the latter regime having the ability to separately fine up to the greater of £17.5 million or 4% of global turnover. The relationship between the United Kingdom and the EU in relation to certain aspects of data protection law remains unclear, for example around how data can lawfully be transferred between each jurisdiction, which may expose us to further compliance risk. The European Commission has adopted an adequacy decision in favor of the UK, enabling data transfers from EU member states to the UK without additional safeguards. However, the UK adequacy decision will automatically expire in June 2025 unless the European Commission re-assesses and renews/ extends that decision.

We are also subject to evolving EU and EEA privacy laws on cookies and e-marketing. The EU is in the process of replacing the e-Privacy Directive (2002/58/EC) with a new set of rules taking the form of a regulation, which will be directly applicable in each European Member State. The draft e-Privacy Regulation imposes strict opt-in marketing rules with limited exceptions for business-to-business communications, alters rules on third-party cookies, web beacons and similar technology and significantly increases fining powers to the same levels as the GDPR (i.e. the greater of EUR 20 million or 4% of total global annual turnover for certain breaches). We have enlisted the help of external advisors to provide assistance in implementing a robust program that we believe will ensure compliance with these obligations which is likely to require us to expend capital and other resources.

Any actual or perceived failure by us, our employees or contractors, our partners, our service providers, or the third parties with whom we work, to comply with privacy or security laws, policies, legal obligations or industry standards, or any security incident that results in the unauthorized release or transfer of personal information, may result in governmental enforcement actions and investigations including by EU regulators and U.S. federal and state regulatory authorities as well as fines and penalties, litigation, including by consumer advocacy groups, and/or adverse publicity and could cause our customers, their patients and other healthcare professionals to lose trust in us, which could harm our reputation and have a material adverse effect on our business, financial condition and results of operations.

***Healthcare policy changes, including recently enacted legislation reforming the U.S. healthcare system, could harm our business, financial condition and results of operations.***

In the U.S., there have been and continue to be a number of legislative initiatives to contain healthcare costs. In March 2010, the Affordable Care Act was enacted in the U.S., which made a number of substantial changes in the way healthcare is financed by both governmental and private insurers. Among other ways in which it may affect our business, the Affordable Care Act:

- established a new Patient-Centered Outcomes Research Institute to oversee and identify priorities in comparative clinical effectiveness research in an effort to coordinate and develop such research;
- implemented payment system reforms including a national pilot program on payment bundling to encourage hospitals, physicians and other providers to improve the coordination, quality and efficiency of certain healthcare services through bundled payment models; and
- expanded the eligibility criteria for Medicaid programs.

The expansion in the government's role in the U.S. healthcare industry may result in decreased profits to us, lower reimbursement by payors for our Inspire system, and/or reduced medical procedure volumes, all of which may have a material adverse effect on our business, financial condition and results of operations.

Since its enactment, there have been judicial, executive and Congressional challenges to the Affordable Care Act. On June 17, 2021, the U.S. Supreme Court dismissed the most recent judicial challenge to the Affordable Care Act without specifically ruling on the constitutionality of the Affordable Care Act. Prior to the Supreme Court's decision, President Biden issued an executive order to initiate a special enrollment period from February 15, 2021 through August 15, 2021 for purposes of obtaining health insurance coverage through the Affordable Care Act marketplace. The executive order also instructed certain governmental agencies to review and reconsider their existing policies and rules that limit access to healthcare, including among others, reexamining Medicaid demonstration projects and waiver programs that include work requirements, and policies that create unnecessary barriers to obtaining access to health insurance coverage through Medicaid or the Affordable Care Act.

In addition, other legislative changes have been proposed and adopted since the Affordable Care Act was enacted. On August 2, 2011, the Budget Control Act of 2011 was signed into law, which, among other things, reduced Medicare payments to providers by 2% per fiscal year, effective on April 1, 2013 and, due to subsequent legislative amendments to the statute, will remain in effect through 2030, with the exception of a temporary suspension from May 1, 2020 through March 31, 2022, unless additional Congressional action is taken. On January 2, 2013, the American Taxpayer Relief Act of 2012 was signed into law, which, among other things, reduced Medicare payments to several providers, including hospitals, and increased the statute of limitations period for the government to recover overpayments to providers from three to five years. In addition, the Medicare Access and CHIP Reauthorization Act of 2015 ("MACRA") enacted on April 16, 2015, repealed the formula by which Medicare made annual payment adjustments to physicians and replaced the former formula with fixed annual updates and a new system of incentive payments began in 2019 that are based on various performance measures and physicians' participation in alternative payment models such as accountable care organizations. It is unclear what effect new quality and payment programs, such as MACRA, may have on our business, financial condition, results of operations or cash flows.

We expect additional state, federal, and foreign healthcare policies and reform measures to be adopted in the future, any of which could limit reimbursement for healthcare products and services or otherwise result in reduced demand for our Inspire system or additional pricing pressure and have a material adverse effect on our industry generally and on our customers. Any changes of, or uncertainty with respect to, future coverage or reimbursement rates could affect demand for our Inspire system, which in turn could impact our ability to successfully commercialize our Inspire system and could have a material adverse effect on our business, financial condition and results of operations.

***Our business involves the use of hazardous materials and our third-party manufacturers must comply with environmental laws and regulations, which may be expensive and restrict how we do business.***

Our third-party manufacturers' activities may involve the controlled storage, use and disposal of hazardous materials. Our manufacturers are subject to federal, state, local, and foreign laws and regulations governing the use, generation, manufacture, storage, handling and disposal of these hazardous materials. We currently carry no insurance specifically covering environmental claims relating to the use of hazardous materials. Although we believe the safety procedures of our manufacturers for handling and disposing of these materials and waste products comply with the standards prescribed by these laws and regulations, we cannot eliminate the risk of accidental injury or contamination from the use, storage, handling or disposal of hazardous materials. In the event



of an accident, state or federal or other applicable authorities may curtail our manufacturers' use of these materials and interrupt their business operations which could adversely affect our business.

***The clinical trial process required to obtain regulatory approvals is lengthy and expensive with uncertain outcomes. If clinical studies of our future products do not produce results necessary to support regulatory clearance or approval in the U.S. or foreign approval or certification, with respect to our current or future products, elsewhere, we will be unable to expand the indications for or commercialize these products and may incur additional costs or experience delays in completing, or ultimately be unable to complete, the commercialization of those products.***

We have obtained PMA for our Inspire system. In order to obtain PMA for a device, the sponsor must conduct well-controlled clinical trials designed to assess the safety and efficacy of the product candidate. Conducting clinical trials is a complex and expensive process, can take many years, and outcomes are inherently uncertain. We incur substantial expense for, and devote significant time to, clinical trials but cannot be certain that the trials will ever result in commercial revenue. We may experience significant setbacks in clinical trials, even after earlier clinical trials showed promising results, and failure can occur at any time during the clinical trial process. Any of our products may malfunction or may produce undesirable adverse effects that could cause us or regulatory authorities to interrupt, delay or halt clinical trials. We, the FDA, or another regulatory authority may suspend or terminate clinical trials at any time to avoid exposing trial participants to unacceptable health risks.

Successful results of pre-clinical studies are not necessarily indicative of future clinical trial results, and predecessor clinical trial results may not be replicated in subsequent clinical trials. Additionally, the FDA or foreign regulatory authorities or notified bodies may disagree with our interpretation of the data from our pre-clinical studies and clinical trials, or may find the clinical trial design, conduct or results inadequate to prove safety or efficacy, and may require us to pursue additional pre-clinical studies or clinical trials, which could further delay the clearance, approval or certification of our products. The data we collect from our pre-clinical studies and clinical trials may not be sufficient to support FDA clearance or approval or foreign approval or certification, and if we are unable to demonstrate the safety and efficacy of our future products in our clinical trials, we will be unable to obtain regulatory clearance, approval or certification to market our products.

In addition, we may estimate and publicly announce the anticipated timing of the accomplishment of various clinical, regulatory and other product development goals, which are often referred to as milestones. These milestones could include the obtainment of the right to affix the CE mark in the EU; the submission to the FDA of an IDE application to commence a pivotal clinical trial for a new product candidate; the enrollment of patients in clinical trials; the release of data from clinical trials; and other clinical and regulatory events. The actual timing of these milestones could vary dramatically compared to our estimates, in some cases for reasons beyond our control. We cannot ensure that we will meet our projected milestones and if we do not meet these milestones as publicly announced, the commercialization of our products may be delayed and, as a result, our stock price may decline.

Clinical trials are necessary to support PMA applications and may be necessary to support PMA supplements for modified versions of our marketed device products. This would require the enrollment of large numbers of suitable subjects, which may be difficult to identify, recruit and maintain as participants in the clinical trial. The clinical trials supporting the PMA application for our Inspire system involved 126 randomized patients. Adverse outcomes in the post-approval studies could also result in restrictions or withdrawal of approval of the PMA. We will likely need to conduct additional clinical studies in the future to support new indications for our products or for approvals, clearances or certifications of new product lines, or for the approval of the use of our products in some foreign countries. Clinical testing is difficult to design and implement, can take many years, can be expensive and carries uncertain outcomes. The initiation and completion of any of these studies may be prevented, delayed, or halted for numerous reasons. We may experience a number of events during that could adversely affect the costs, timing or successful completion of our clinical trials, including:

- we may be required to submit an IDE application or similar application to the FDA or a foreign regulatory authority, which must become effective prior to commencing human clinical trials, and the FDA or foreign regulatory authority may reject our IDE or similar application and notify us that we may not begin investigational trials;

- regulators and other comparable foreign regulatory authorities may disagree as to the design or implementation of our clinical trials;
- regulators and/or IRBs, or other reviewing bodies may not authorize us or our investigators to commence a clinical trial, or to conduct or continue a clinical trial at a prospective or specific trial site;
- we may not reach agreement on acceptable terms with prospective contract research organizations ("CROs") and clinical trial sites, the terms of which can be subject to extensive negotiation and may vary significantly among different CROs and trial sites;
- clinical trials may produce negative or inconclusive results, and we may decide, or regulators may require us, to conduct additional clinical trials or abandon product development programs;
- the number of subjects or patients required for clinical trials may be larger than we anticipate, enrollment in these clinical trials may be insufficient or slower than we anticipate, and the number of clinical trials being conducted at any given time may be high and result in fewer available patients for any given clinical trial, or patients may drop out of these clinical trials at a higher rate than we anticipate;
- our third-party contractors, including those manufacturing products or conducting clinical trials on our behalf, may fail to comply with regulatory requirements or meet their contractual obligations to us in a timely manner, or at all;
- we might have to suspend or terminate clinical trials for various reasons, including a finding that the subjects are being exposed to unacceptable health risks;
- we may have to amend clinical trial protocols or conduct additional studies to reflect changes in regulatory requirements or guidance, which we may be required to submit to an IRB or other review bodies and/or regulatory authorities for re-examination;
- regulators, IRBs, or other parties may require or recommend that we or our investigators suspend or terminate clinical research for various reasons, including safety signals or noncompliance with regulatory requirements;
- the cost of clinical trials may be greater than we anticipate;
- clinical sites may not adhere to the clinical protocol or may drop out of a clinical trial;
- we may be unable to recruit a sufficient number of clinical trial sites;
- regulators, IRBs, or other reviewing bodies may fail to approve or subsequently find fault with our manufacturing processes or facilities of third-party manufacturers with which we enter into agreement for clinical and commercial supplies, the supply of devices or other materials necessary to conduct clinical trials may be insufficient, inadequate or not available at an acceptable cost, or we may experience interruptions in supply;
- approval policies or regulations of FDA or applicable foreign regulatory agencies may change in a manner rendering our clinical data insufficient for approval or certification; and
- our current or future products may have undesirable side effects or other unexpected characteristics.

Patient enrollment in clinical trials and completion of patient follow-up depend on many factors, including the size of the patient population, the nature of the trial protocol, the proximity of patients to clinical sites, the eligibility criteria for the clinical trial, patient compliance, competing clinical trials and clinicians' and patients' perceptions as to the potential advantages of the product being studied in relation to other available therapies, including any new treatments that may be approved for the indications we are investigating. For example, patients may be discouraged from enrolling in our clinical trials if the trial protocol requires them to undergo extensive post-

treatment procedures or follow-up to assess the safety and efficacy of a product candidate, or they may be persuaded to participate in contemporaneous clinical trials of a competitor's product candidate. In addition, patients participating in our clinical trials may drop out before completion of the trial or experience adverse medical events unrelated to our products. Delays in patient enrollment or failure of patients to continue to participate in a clinical trial may delay commencement or completion of the clinical trial, cause an increase in the costs of the clinical trial and delays, or result in the failure of the clinical trial.

Clinical trials must be conducted in accordance with the laws and regulations of the FDA and other applicable regulatory authorities' legal requirements, regulations or guidelines, and are subject to oversight by these governmental agencies and IRBs and other reviewing bodies at the medical institutions where the clinical trials are conducted. In addition, clinical trials must be conducted with supplies of our devices produced under current good manufacturing practice requirements and other regulations. Furthermore, we rely on CROs, and clinical trial sites to ensure the proper and timely conduct of our clinical trials and while we have agreements governing their committed activities, we have limited influence over their actual performance. We depend on our collaborators and on medical institutions and CROs to conduct our clinical trials in compliance with good clinical practice ("GCP") requirements. To the extent our collaborators or the CROs fail to enroll participants for our clinical trials, fail to conduct the study in accordance with GCP requirements or are delayed for a significant time in the execution of trials, including achieving full enrollment, we may be affected by increased costs, program delays or both. In addition, clinical trials that are conducted in countries outside the U.S. may subject us to further delays and expenses as a result of increased shipment costs, additional regulatory requirements and the engagement of non-U.S. CROs, as well as expose us to risks associated with clinical investigators who are unknown to the FDA, and different standards of diagnosis, screening and medical care.

Failure can occur at any stage of clinical testing. Our clinical studies may produce negative or inconclusive results, and we may decide, or regulators may require us, to conduct additional clinical and non-clinical testing in addition to those we have planned. Our failure to adequately demonstrate the safety and efficacy of our system or any product we may develop in the future would prevent receipt of regulatory clearance, approval or certification and, ultimately, the commercialization of that product or indication for use. Even if our future products are cleared or approved in the U.S., commercialization of our products in foreign countries would require approval by regulatory authorities or certification by notified bodies in those countries. Approval and certification procedures vary among jurisdictions and can involve requirements and administrative review periods different from, and greater than, those in the U.S., including additional preclinical studies or clinical trials. Any of these occurrences could have an adverse effect on our business, financial condition and results of operations.

***Disruptions at the FDA, other government agencies or notified bodies caused by funding shortages or global health concerns could hinder their ability to hire and retain key leadership and other personnel, or otherwise prevent new products and services from being developed or commercialized in a timely manner, which could negatively impact our business.***

The ability of the FDA, foreign regulatory authorities and notified bodies to review and approve or certify new products can be affected by a variety of factors, including government budget and funding levels, ability to hire and retain key personnel and accept the payment of user fees, and statutory, regulatory, and policy changes. Average review times at the agency have fluctuated in recent years as a result. In addition, government funding of other government agencies that fund research and development activities is subject to the political process, which is inherently fluid and unpredictable.

Disruptions at the FDA and other agencies or notified bodies may also slow the time necessary for new medical devices and modifications to cleared or approved medical devices to be reviewed and/or cleared, approved or certified by necessary government agencies or notified bodies, which would adversely affect our business. For example, over the last several years, the U.S. government has shut down several times and certain regulatory agencies, such as the FDA, have had to furlough critical FDA employees and stop critical activities. If a prolonged government shutdown occurs, it could significantly impact the ability of the FDA to timely review and process our regulatory submissions, which could have a material adverse effect on our business. Similarly, a prolonged government shutdown could prevent the timely review of our patent applications by the United States Patent and Trademark Office ("USPTO"), which could delay the issuance of any U.S. patents to which we might otherwise be

entitled. Further, in our operations as a public company, future government shutdowns could impact our ability to access the public markets and obtain necessary capital in order to properly fund our business.

Separately, in response to the COVID-19 pandemic, in March 2020, the FDA announced its intention to postpone most inspections of foreign manufacturing facilities, and on March 18, 2020, the FDA temporarily postponed routine surveillance inspections of domestic manufacturing facilities. Subsequently, in July 2020, the FDA resumed certain on-site inspections of domestic manufacturing facilities subject to a risk-based prioritization system. The FDA utilized this risk-based assessment system to assist in determining when and where it was safest to conduct prioritized domestic inspections. In May 2021, the FDA outlined a detailed plan to move toward a more consistent state of inspectional operations, and in July 2021, the FDA resumed standard inspectional operations of domestic facilities and was continuing to maintain this level of operation as of September 2021. More recently, the FDA has continued to monitor and implement changes to its inspectional activities to ensure the safety of its employees and those of the firms it regulates as it adapts to the evolving COVID-19 pandemic.

In the European Union ("EU"), notified bodies must be officially designated to certify products and services in accordance with the EU Medical Devices Regulation. While several notified bodies have been designated, the COVID-19 pandemic has significantly slowed down the designation process. Currently designated notified bodies have severe capacity constraints and are facing a large amount of requests for recertification of products under the MDR as a consequence of which review times have lengthened. This situation could significantly impact our ability to grow our business in the EU and EEA.

Regulatory authorities outside the U.S. may adopt similar restrictions or other policy measures in response to the COVID-19 pandemic. If a prolonged government shutdown occurs, or if global health concerns continue to prevent the FDA, other regulatory authorities and notified bodies from conducting their regular inspections or audits, reviews, or other regulatory activities, it could significantly impact the ability of the FDA, other regulatory authorities or notified bodies to timely review and process our regulatory submissions, which could have a material adverse effect on our business.

## **Risks Related to Intellectual Property Matters**

***If we are unable to adequately protect our intellectual property rights, or if we are accused of infringing on the intellectual property rights of others, our competitive position could be harmed or we could be required to incur significant expenses to enforce or defend our rights.***

Our commercial success will depend in part on our success in obtaining and maintaining issued patents, trademarks and other intellectual property rights in the U.S. and elsewhere and protecting our proprietary technology. If we do not adequately protect our intellectual property and proprietary technology, competitors may be able to use our technologies or the goodwill we have acquired in the marketplace and erode or negate any competitive advantage we may have, which could harm our business and ability to achieve profitability.

Some of our intellectual property rights depend on a licensing agreement with a third party, and our patent coverage includes protection provided by licensed patents. Many of these licensed patents are over ten years old and the standard life of a patent is 20 years from its initial filing date. If in the future we no longer have rights to one or more of these licensed patents, our patent coverage may be compromised, which in turn could affect our ability to protect our Inspire system or defend against competitors.

We own numerous issued patents and pending patent applications that relate to our system. As of December 31, 2021, we had rights to 44 issued U.S. patents, 34 issued foreign patents, 75 pending U.S. patent applications, and 63 pending foreign patent applications. Assuming all required fees are paid, issued U.S. patents owned by us will expire between 2022 and 2037.

We cannot provide any assurances that any of our patents have, or that any of our pending patent applications that mature into issued patents will include, claims with a scope sufficient to protect our Inspire system, any additional features we develop for our Inspire system or any new products. Other parties may have developed technologies that may be related or competitive to our system, may have filed or may file patent applications and

may have received or may receive patents that overlap or conflict with our patent applications, either by claiming the same methods or devices or by claiming subject matter that could dominate our patent position. The patent positions of medical device companies, including our patent position, may involve complex legal and factual questions, and, therefore, the scope, validity and enforceability of any patent claims that we may obtain cannot be predicted with certainty. Patents, if issued, may be challenged, deemed unenforceable, invalidated or circumvented. Proceedings challenging our patents could result in either loss of the patent or denial of the patent application or loss or reduction in the scope of one or more of the claims of the patent or patent application. In addition, such proceedings may be costly. Thus, any patents that we may own may not provide any protection against competitors. Furthermore, an adverse decision in an interference proceeding can result in a third party receiving the patent right sought by us, which in turn could affect our ability to commercialize our products.

Though an issued patent is presumed valid and enforceable, its issuance is not conclusive as to its validity or its enforceability and it may not provide us with adequate proprietary protection or competitive advantages against competitors with similar products. Competitors could purchase our Inspire system and attempt to replicate some or all of the competitive advantages we derive from our development efforts, willfully infringe our intellectual property rights, design around our patents, or develop and obtain patent protection for more effective technologies, designs or methods. We may be unable to prevent the unauthorized disclosure or use of our technical knowledge or trade secrets by consultants, suppliers, vendors, former employees and current employees. The laws of some foreign countries do not protect our proprietary rights to the same extent as the laws of the U.S., and we may encounter significant problems in protecting our proprietary rights in these countries.

Our ability to enforce our patent rights depends on our ability to detect infringement. It may be difficult to detect infringers who do not advertise the components that are used in their products. Moreover, it may be difficult or impossible to obtain evidence of infringement in a competitor's or potential competitor's product. We may not prevail in any lawsuits that we initiate and the damages or other remedies awarded if we were to prevail may not be commercially meaningful.

In addition, proceedings to enforce or defend our patents could put our patents at risk of being invalidated, held unenforceable or interpreted narrowly. Such proceedings could also provoke third parties to assert claims against us, including that some or all of the claims in one or more of our patents are invalid or otherwise unenforceable. If any of our patents covering our Inspire system are invalidated or found unenforceable, or if a court found that valid, enforceable patents held by third parties covered one or more of our products, our competitive position could be harmed or we could be required to incur significant expenses to enforce or defend our rights.

The degree of future protection for our proprietary rights is uncertain, and we cannot ensure that:

- any of our patents, or any of our pending patent applications, if issued, will include claims having a scope sufficient to protect our Inspire system;
- any of our pending patent applications will issue as patents;
- we will be able to successfully commercialize our products on a substantial scale, if approved, before our relevant patents we may have expire;
- we were the first to make the inventions covered by each of our patents and pending patent applications;
- we were the first to file patent applications for these inventions;
- others will not develop similar or alternative technologies that do not infringe our patents; any of our patents will be found to ultimately be valid and enforceable;
- any patents issued to us will provide a basis for an exclusive market for our commercially viable products, will provide us with any competitive advantages or will not be challenged by third parties;
- we will develop additional proprietary technologies or products that are separately patentable; or

- our commercial activities or products will not infringe upon the patents of others.

We rely, in part, upon unpatented trade secrets, unpatented know-how and continuing technological innovation to develop and maintain our competitive position, which we seek to protect, in part, by confidentiality agreements with our employees and our collaborators and consultants. We also have agreements with our employees and consultants that obligate them to assign their inventions to us and have non-compete agreements with some, but not all, of our consultants. It is possible that technology relevant to our business will be independently developed by a person that is not a party to such an agreement. Furthermore, if the employees and consultants who are parties to these agreements breach or violate the terms of these agreements, we may not have adequate remedies for any such breach or violation, and we could lose our trade secrets through such breaches or violations. Further, our trade secrets could otherwise become known or be independently discovered by our competitors.

***Litigation or other proceedings or third-party claims of intellectual property infringement could require us to spend significant time and money and could prevent us from selling our products or affect our stock price.***

Our commercial success will depend in part on not infringing the patents or violating the other proprietary rights of others. Significant litigation regarding patent rights occurs in our industry. Our competitors in both the U.S. and abroad, many of which have substantially greater resources and have made substantial investments in patent portfolios and competing technologies, may have applied for or obtained or may in the future apply for and obtain, patents that will prevent, limit or otherwise interfere with our ability to make, use and sell our products. We do not always conduct independent reviews of patents issued to third parties. In addition, patent applications in the U.S. and elsewhere can be pending for many years before issuance, or unintentionally abandoned patents or applications can be revived, so there may be applications of others now pending or recently revived patents of which we are unaware. These applications may later result in issued patents, or the revival of previously abandoned patents, that will prevent, limit or otherwise interfere with our ability to make, use or sell our products. Third parties may, in the future, assert claims that we are employing their proprietary technology without authorization, including claims from competitors or from non-practicing entities that have no relevant product revenue and against whom our own patent portfolio may have no deterrent effect. As we continue to commercialize our products in their current or updated forms, launch new products and enter new markets, competitors may claim that one or more of our products infringe their intellectual property rights as part of business strategies designed to impede our successful commercialization and entry into new markets. The large number of patents, the rapid rate of new patent applications and issuances, the complexities of the technology involved, and the uncertainty of litigation may increase the risk of business resources and management's attention being diverted to patent litigation. We have, and we may in the future, receive letters or other threats or claims from third parties inviting us to take licenses under, or alleging that we infringe, their patents.

Moreover, we may become party to future adversarial proceedings regarding our patent portfolio or the patents of third parties. Such proceedings could include supplemental examination or contested post-grant proceedings such as review, reexamination, inter partes review, interference or derivation proceedings before the USPTO and challenges in U.S. District Court. Patents may be subjected to opposition, post-grant review or comparable proceedings lodged in various foreign, both national and regional, patent offices. The legal threshold for initiating litigation or contested proceedings may be low, so that even lawsuits or proceedings with a low probability of success might be initiated. Litigation and contested proceedings can also be expensive and time-consuming, and our adversaries in these proceedings may have the ability to dedicate substantially greater resources to prosecuting these legal actions than we can. We may also occasionally use these proceedings to challenge the patent rights of others. We cannot be certain that any particular challenge will be successful in limiting or eliminating the challenged patent rights of the third party.

Any lawsuits resulting from such allegations could subject us to significant liability for damages and invalidate our proprietary rights. Any potential intellectual property litigation also could force us to do one or more of the following:

- stop making, selling or using products or technologies that allegedly infringe the asserted intellectual property;

- lose the opportunity to license our technology to others or to collect royalty payments based upon successful protection and assertion of our intellectual property rights against others; incur significant legal expenses;
- pay substantial damages or royalties to the party whose intellectual property rights we may be found to be infringing;
- pay the attorney's fees and costs of litigation to the party whose intellectual property rights we may be found to be infringing;
- redesign those products that contain the allegedly infringing intellectual property, which could be costly, disruptive and infeasible; and
- attempt to obtain a license to the relevant intellectual property from third parties, which may not be available on reasonable terms or at all, or from third parties who may attempt to license rights that they do not have.

Any litigation or claim against us, even those without merit, may cause us to incur substantial costs, and could place a significant strain on our financial resources, divert the attention of management from our core business and harm our reputation. If we are found to infringe the intellectual property rights of third parties, we could be required to pay substantial damages (which may be increased up to three times of awarded damages) and/or substantial royalties and could be prevented from selling our products unless we obtain a license or are able to redesign our products to avoid infringement. Any such license may not be available on reasonable terms, if at all, and there can be no assurance that we would be able to redesign our products in a way that would not infringe the intellectual property rights of others. We could encounter delays in product introductions while we attempt to develop alternative methods or products. If we fail to obtain any required licenses or make any necessary changes to our products or technologies, we may have to withdraw existing products from the market or may be unable to commercialize one or more of our products.

In addition, we generally indemnify our customers with respect to infringement by our products of the proprietary rights of third parties. Third parties may assert infringement claims against our customers. These claims may require us to initiate or defend protracted and costly litigation on behalf of our customers, regardless of the merits of these claims. If any of these claims succeed or settle, we may be forced to pay damages or settlement payments on behalf of our customers or may be required to obtain licenses for the products they use. If we cannot obtain all necessary licenses on commercially reasonable terms, our customers may be forced to stop using our products.

***If we are unable to protect the confidentiality of our trade secrets, our business and competitive position could be harmed.***

In addition to patent protection for our issued patents and pending patent applications related to our system, we also rely upon copyright and trade secret protection for our Inspire therapy, as well as non-disclosure agreements and invention assignment agreements with our employees, consultants and third parties, to protect our confidential and proprietary information. In addition to contractual measures, we try to protect the confidential nature of our proprietary information using commonly accepted physical and technological security measures. Such measures may not, for example, in the case of misappropriation of a trade secret by an employee or third party with authorized access, provide adequate protection for our proprietary information. Our security measures may not prevent an employee or consultant from misappropriating our trade secrets and providing them to a competitor, and recourse we take against such misconduct may not provide an adequate remedy to protect our interests fully. Unauthorized parties may also attempt to copy or reverse engineer certain aspects of our products that we consider proprietary. Enforcing a claim that a party illegally disclosed or misappropriated a trade secret can be difficult, expensive and time-consuming, and the outcome is unpredictable. Even though we use commonly accepted security measures, trade secret violations are often a matter of state law, and the criteria for protection of trade secrets can vary among different jurisdictions. In addition, trade secrets may be independently developed by others in a manner that could prevent legal recourse by us. If any of our confidential or proprietary

information, such as our trade secrets, were to be disclosed or misappropriated, or if any such information was independently developed by a competitor, our business and competitive position could be harmed.

***We may be unable to enforce our intellectual property rights throughout the world.***

The laws of some foreign countries do not protect intellectual property rights to the same extent as the laws of the U.S. Many companies have encountered significant problems in protecting and defending intellectual property rights in certain foreign jurisdictions. This could make it difficult for us to stop infringement of our foreign patents, if obtained, or the misappropriation of our other intellectual property rights. For example, some foreign countries have compulsory licensing laws under which a patent owner must grant licenses to third parties. In addition, some countries limit the enforceability of patents against third parties, including government agencies or government contractors. In these countries, patents may provide limited or no benefit. Patent protection must ultimately be sought on a country-by-country basis, which is an expensive and time-consuming process with uncertain outcomes. Accordingly, we may choose not to seek patent protection in certain countries, and we will not have the benefit of patent protection in such countries.

Proceedings to enforce our patent rights in foreign jurisdictions could result in substantial costs and divert our efforts and attention from other aspects of our business. Accordingly, our efforts to protect our intellectual property rights in such countries may be inadequate. In addition, changes in the law and legal decisions by courts in the U.S. and foreign countries may affect our ability to obtain adequate protection for our technology and the enforcement of our intellectual property.

***Third parties may assert ownership or commercial rights to inventions we develop.***

Third parties may in the future make claims challenging the inventorship or ownership of our intellectual property. We have written agreements with collaborators that provide for the ownership of intellectual property arising from our collaborations. In addition, we may face claims by third parties that our agreements with employees, contractors or consultants obligating them to assign intellectual property to us are ineffective or in conflict with prior or competing contractual obligations of assignment, which could result in ownership disputes regarding intellectual property we have developed or will develop and interfere with our ability to capture the commercial value of such intellectual property. Litigation may be necessary to resolve an ownership dispute, and if we are not successful, we may be precluded from using certain intellectual property or may lose our exclusive rights in that intellectual property. Either outcome could harm our business and competitive position.

***Third parties may assert that our employees or consultants have wrongfully used or disclosed confidential information or misappropriated trade secrets.***

We employ individuals who previously worked with other companies, including our competitors or potential competitors. Although we try to ensure that our employees and consultants do not use the proprietary information or know-how of others in their work for us, we may be subject to claims that we or our employees, consultants or independent contractors have inadvertently or otherwise used or disclosed intellectual property or personal data, including trade secrets or other proprietary information, of a former employer or other third party. Litigation may be necessary to defend against these claims. If we fail in defending any such claims or settling those claims, in addition to paying monetary damages or a settlement payment, we may lose valuable intellectual property rights or personnel. Even if we are successful in defending against such claims, litigation could result in substantial costs and be a distraction to management and other employees.

***Recent changes in U.S. patent laws may limit our ability to obtain, defend and/or enforce our patents.***

Recent patent reform legislation could increase the uncertainties and costs surrounding the prosecution of our patent applications and the enforcement or defense of our issued patents. The Leahy-Smith America Invents Act ("the Leahy-Smith Act") includes a number of significant changes to U.S. patent law. These include provisions that affect the way patent applications are prosecuted and also affect patent litigation. The USPTO recently developed new regulations and procedures to govern administration of the Leahy-Smith Act, and many of the substantive changes to patent law associated with the Leahy-Smith Act, and in particular, the first to file provisions, which became effective on March 16, 2013. The first to file provisions limit the rights of an inventor to



patent an invention if not the first to file an application for patenting that invention, even if such invention was the first invention. Accordingly, it is not clear what, if any, impact the Leahy-Smith Act will have on the operation of our business.

However, the Leahy-Smith Act and its implementation could increase the uncertainties and costs surrounding the enforcement and defense of our issued patents. For example, the Leahy-Smith Act provides that an administrative tribunal known as the Patent Trial and Appeals Board ("PTAB") provides a venue for challenging the validity of patents at a cost that is much lower than district court litigation and on timelines that are much faster. Although it is not clear what, if any, long-term impact the PTAB proceedings will have on the operation of our business, the initial results of patent challenge proceedings before the PTAB since its inception in 2013 have resulted in the invalidation of many U.S. patent claims. The availability of the PTAB as a lower-cost, faster and potentially more potent tribunal for challenging patents could increase the likelihood that our own patents will be challenged, thereby increasing the uncertainties and costs of maintaining and enforcing them.

## **Risks Related to Our Common Stock**

### ***The market price of our common stock may be volatile and fluctuate substantially, which could result in substantial losses for purchasers of our common stock***

The market price of our common stock is likely to be highly volatile and may fluctuate substantially due to many factors, including:

- the volume and timing of sales of our products;
- the introduction of new products or product enhancements by us or others in our industry;
- disputes or other developments with respect to our or others' intellectual property rights;
- our ability to develop, obtain regulatory clearance or approval for, and market new and enhanced products on a timely basis;
- regulatory actions with respect to our therapy or those of our competitors or companies perceived to be similar to ours;
- product liability claims or other litigation;
- changes in physician, hospital, healthcare provider practices;
- quarterly variations in our results of operations or those of others in our industry;
- media exposure of our products or of those of others in our industry;
- changes in governmental regulations
- changes in the structure of healthcare payment systems;
- changes in earnings estimates or recommendations by securities analysts; and
- general market conditions and other factors, including factors unrelated to our operating performance or the operating performance of our competitors.

In recent years, the stock markets generally have experienced extreme price and volume fluctuations that have often been unrelated or disproportionate to the operating performance of those companies. Broad market and industry factors may significantly affect the market price of our common stock, regardless of our actual operating performance.

In addition, in the past, class action litigation has often been instituted against companies whose securities have experienced periods of volatility in market price. Securities litigation brought against us following volatility in our stock price, regardless of the merit or ultimate results of such litigation, could result in substantial costs, which would hurt our financial condition and operating results and divert management's attention and resources from our business.

***Provisions in our governing documents and under Delaware law could make an acquisition of us more difficult and may prevent attempts by our stockholders to replace or remove our current management.***

Provisions in our amended and restated certificate of incorporation and our amended and restated bylaws may discourage, delay or prevent a merger, acquisition or other change in control of us that stockholders may consider favorable, including transactions in which stockholders might otherwise receive a premium for their shares. These provisions could also 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. In addition, these provisions may frustrate or prevent any attempts by our stockholders to replace or remove our current management by making it more difficult for stockholders to replace members of our board of directors. Because our board of directors is responsible for appointing the members of our management team, these provisions could in turn affect any attempt by our stockholders to replace current members of our management team. These provisions provide, among other things, that:

- our board of directors has the exclusive right to expand the size of our board of directors and to elect directors to fill a vacancy created by the expansion of the board of directors or the resignation, death or removal of a director, which prevents stockholders from being able to fill vacancies on our board of directors;
- our board of directors is divided into three classes, Class I, Class II and Class III, with each class serving staggered three-year terms, which may delay the ability of stockholders to change the membership of a majority of our board of directors;
- our stockholders may not act by written consent, which forces stockholder action to be taken at an annual or special meeting of our stockholders;
- a special meeting of stockholders may be called only by the chair of our board of directors, our chief executive officer or a majority of our board of directors, which may delay the ability of our stockholders to force consideration of a proposal or to take action, including the removal of directors;
- our amended and restated certificate of incorporation prohibits cumulative voting in the election of directors, which limits the ability of minority stockholders to elect director candidates;
- our board of directors may alter certain provisions of our bylaws without obtaining stockholder approval;
- the approval of the holders of at least two-thirds of the shares entitled to vote at an election of directors is required to adopt, amend or repeal our bylaws or repeal the provisions of our amended and restated certificate of incorporation regarding the election and removal of directors;
- stockholders must provide advance notice and additional disclosures in order to nominate individuals for election to the board of directors or to propose matters that can be acted upon at a stockholders' meeting, which may discourage or deter a potential acquiror from conducting a solicitation of proxies to elect the acquiror's own slate of directors or otherwise attempting to obtain control of our Company; and
- our board of directors is authorized to issue shares of preferred stock and to determine the terms of those shares, including preferences and voting rights, without stockholder approval, which could be used to significantly dilute the ownership of a hostile acquiror.

Moreover, because we are incorporated in Delaware, we are governed by the provisions of Section 203 of the Delaware General Corporation Law, which prohibits a person who owns in excess of 15% of our outstanding

voting stock from merging or combining with us for a period of three years after the date of the transaction in which the person acquired in excess of 15% of our outstanding voting stock, unless the merger or combination is approved in a prescribed manner.

***Our amended and restated certificate of incorporation provides that the Court of Chancery of the State of Delaware will be the exclusive forum for substantially all disputes between us and our stockholders, which could limit our stockholders' ability to obtain a favorable judicial forum for disputes with us or our directors, officers or employees.***

Our amended and restated certificate of incorporation provides that the Court of Chancery of the State of Delaware is the exclusive forum for (i) any derivative action or proceeding brought on our behalf, (ii) any action asserting a claim of breach of a fiduciary duty or other wrongdoing by any of our directors, officers, employees or agents to us or our stockholders, (iii) any action asserting a claim arising pursuant to any provision of the DGCL or our amended and restated certificate of incorporation or amended and restated bylaws or (iv) any action asserting a claim governed by the internal affairs doctrine. We believe this provision benefits us by providing increased consistency in the application of Delaware law by chancellors particularly experienced in resolving corporate disputes, efficient administration of cases on a more expedited schedule relative to other forums and protection against the burdens of multi-forum litigation. However, this provision may limit a stockholder's ability to bring a claim in a judicial forum that it finds favorable for disputes with us or our directors, officers or other employees, which may discourage such lawsuits against us and our directors, officers and other employees. Alternatively, if a court were to find the choice of forum provision contained in our amended and restated certificate of incorporation to be inapplicable or unenforceable in an action, we may incur additional costs associated with resolving such action in other jurisdictions.

***Because we do not anticipate paying any cash dividends on our capital stock in the foreseeable future, capital appreciation, if any, will be your sole source of gain.***

We have never declared or paid cash dividends on our capital stock. We currently intend to retain all of our future earnings, if any, to finance the growth and development of our business. In addition, the agreement governing our credit facility precludes, and any future debt agreements may preclude, us from paying cash dividends. As a result, capital appreciation, if any, of our common stock will be your sole source of gain for the foreseeable future.

## **General Risk Factors**

***Our disclosure controls and procedures may not prevent or detect all errors or acts of fraud.***

We have designed our disclosure controls and procedures to provide reasonable assurance that information we must disclose in reports we file or submit under the Exchange Act is accumulated and communicated to management, and recorded, processed, summarized and reported within the time periods specified in the rules and forms of the SEC. We believe that any disclosure controls and procedures, no matter how well-conceived and operated, can provide only reasonable, not absolute, assurance that the objectives of the control system are met.

These inherent limitations include the realities that judgments in decision-making can be faulty, and that breakdowns can occur because of simple error or mistake. Additionally, controls can be circumvented by the individual acts of some persons, by collusion of two or more people or by an unauthorized override of the controls. Accordingly, because of the inherent limitations in our control system, misstatements due to error or fraud may occur and not be detected.

***Securities analysts may not publish favorable research or reports about our business or may publish no information at all, which could cause our stock price or trading volume to decline.***

The trading market for our common stock relies in part on the research and reports that securities or industry analysts publish about us or our business. We do not control these analysts. If any of the analysts who cover us provide inaccurate or unfavorable research or issue an adverse opinion regarding our stock price, our stock price could decline. If one or more of the analysts covering our business downgrade our stock or change their opinion

of our stock, our stock price would likely decline. In addition, if one or more of these analysts cease coverage of our company or fail to regularly publish reports on us, we could lose visibility in the financial markets, which could cause our share price or trading volume to decline.

**Item 1B. Unresolved Staff Comments.**

None.

**Item 2. Properties.**

Our principal office is located in Golden Valley, Minnesota, where we lease approximately 70,000 square feet of office space. We lease this space under non-cancelable operating lease agreements that expire May 31, 2028, with options to renew for one additional period of five years. We intend to add new facilities as we grow and we believe that suitable additional or substitute space will be available as needed to accommodate any such expansion of our operations.

**Item 3. Legal Proceedings.**

From time to time we may be involved in claims and proceedings arising in the course of our business. The outcome of any such claims or proceedings, regardless of the merits, is inherently uncertain. We are not party to any material legal proceedings.

**Item 4. Mine Safety Disclosures.**

Not applicable.

**PART II**

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

**Market Information**

Our common stock trades on the NYSE under the symbol "INSP."

**Holders**

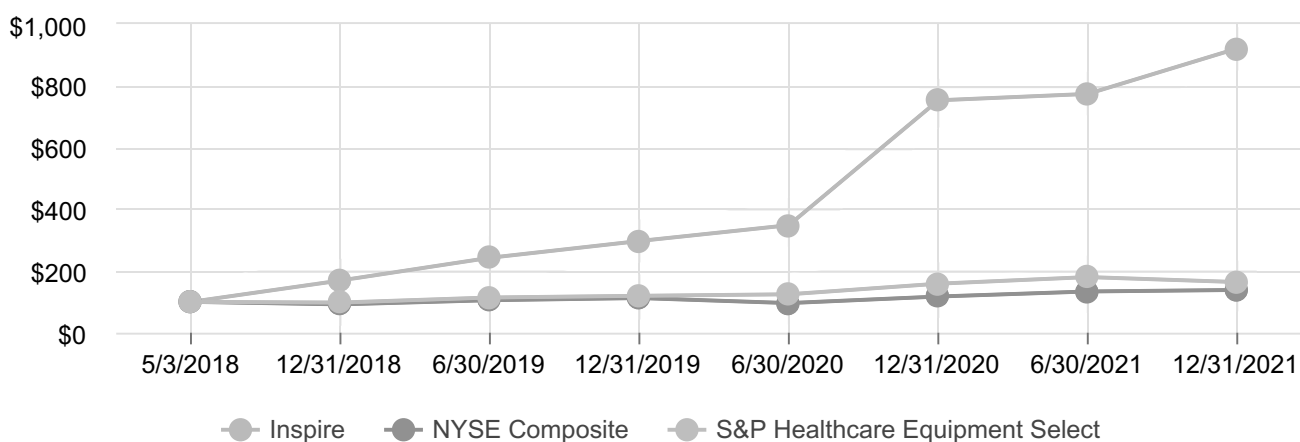
As of February 7, 2022, there were approximately 17 holders of record of our common stock. This number does not include stockholders who are beneficial owners, but whose shares are held in street name by brokers and other nominees. This number of holders of record also does not include stockholders whose shares may be held in trust by other entities.

**Recent Sales of Unregistered Securities**

None.

## Performance Graph

The following graph illustrates a comparison of the total cumulative stockholder return on our common stock with the total return for (i) the S&P Healthcare Equipment Select Industry Index and (ii) the NYSE Composite for the period from May 3, 2018 (the date our common stock commenced trading on the NYSE) through December 31, 2021. The graph assumes an investment of \$100 in our common stock at market close on May 3, 2018 and the reinvestment of dividends, if any. The comparisons in the table are not intended to forecast or be indicative of possible future performance of our common stock. This graph shall not be deemed "soliciting material" or be deemed "filed" for purposes of Section 18 of the Exchange Act, or otherwise subject to the liabilities under that Section, and shall not be deemed to be incorporated by reference into any of our filings under the Securities Act of 1933, as amended (the "Securities Act"), whether made before or after the date hereof and irrespective of any general incorporation language in any such filing.



Stock or Index	Ticker	2018		2019		2020		2021	
		5/3	12/31	6/30	12/31	6/30	12/31	6/30	12/31
Inspire	INSP	\$ 100.00	\$ 169.14	\$ 242.79	\$ 297.08	\$ 348.36	\$ 752.96	\$ 773.66	\$ 920.98
NYSE Composite	NYA	100.00	91.78	105.30	112.27	95.98	117.21	133.59	138.50
S&P Healthcare Equipment Select	SPSIHE	100.00	97.68	114.07	119.57	124.47	158.82	180.23	164.09

## Item 6. [Reserved]

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

*The following discussion and analysis of our financial condition and results of operations should be read in conjunction with our financial statements and the related notes to those statements included elsewhere in this Annual Report on Form 10-K. In addition to historical financial information, the following discussion and analysis contains forward-looking statements that involve risks, uncertainties, and assumptions. Some of the numbers included herein have been rounded for the convenience of presentation. Our actual results may differ materially from those anticipated in these forward-looking statements as a result of many factors, including those discussed under Part I. "Item 1A. Risk Factors" and elsewhere in this Annual Report on Form 10-K.*

### Overview

We are a medical technology company focused on the development and commercialization of innovative, minimally invasive solutions for patients with OSA. Our proprietary Inspire system is the first and only FDA-approved neurostimulation technology that provides a safe and effective treatment for moderate to severe OSA. We have developed a novel, closed-loop solution that continuously monitors a patient's breathing and delivers mild hypoglossal nerve stimulation to maintain an open airway. Inspire therapy is indicated for patients with moderate to severe OSA who do not have significant central sleep apnea and do not have a complete concentric collapse of the airway at the soft palate level. In addition, patients in the U.S. and Japan must have been confirmed to fail or be unable to tolerate positive airway pressure treatments, such as CPAP, and be 18 years of age or older, though there are no similar requirements for patients in Europe.

We sell our Inspire system to hospitals and ASCs in the U.S. and in select countries in Europe through a direct sales organization and we sell our Inspire system in Japan through a distributor. Our direct sales force engages in sales efforts and promotional activities focused on ENT physicians and sleep centers. In addition, we highlight our compelling clinical data and value proposition to increase awareness and adoption amongst referring physicians. We build upon this top-down approach with strong direct-to-consumer marketing initiatives to create awareness of the benefits of our Inspire system and drive interest through patient empowerment. This outreach helps to educate thousands of patients on our Inspire therapy. We increased the number of employees in our sales, marketing, and reimbursement organizations from 47 as of December 31, 2016 to 363 as of December 31, 2021.

Although our sales and marketing efforts are directed at patients and physicians because they are the primary users of our technology, we consider the hospitals and ASCs where the procedure is performed to be our customers, as they are the purchasing agents of our Inspire system. Our customers are reimbursed the cost required to treat each patient through various third-party payors, such as commercial payors and government agencies. Our Inspire system is currently reimbursed primarily on a per-patient prior authorization basis for patients covered by commercial payors, under Local Coverage Determinations for patients covered by Medicare, and under U.S. government contract for patients who are treated by the Veterans Health Administration. As of February 15, 2022, we have secured positive coverage policies with many U.S. commercial payors, including virtually all large national commercial insurers, covering approximately 260 million lives in the U.S. In addition, all seven Medicare Administrative Contractors ("MACs") published final policies in 2020 that provide coverage of Inspire therapy when certain coverage criteria are met.

The procedure performed to implant our device was previously described for billing purposes using a Category I Current Procedural Terminology ("CPT") code (64568), which was used in conjunction with a temporary Category III CPT code (+0466T). At the October 2020 American Medical Association ("AMA") CPT Editorial Panel meeting, the AMA approved the creation of new Category I CPT codes (64582, 64583, and 64584) to separately identify hypoglossal nerve stimulator services. A new Category I code (42975) was also approved for Drug-Induced Sleep Endoscopy, which is the final procedure to determine which patients are appropriate for Inspire therapy. These new codes went into effect on January 1, 2022. With these approvals, a formal survey was conducted to determine the Medicare reimbursement levels assigned to each code and in November 2021 the final 2022 reimbursement rates were announced by the Centers for Medicare and Medicaid Services ("CMS"). The 2022 national average physician payments are approximately \$888 for implantation of a hypoglossal nerve stimulator and approximately \$115 for the DISE procedure. The 2022 rates of Medicare reimbursement to our hospital customers is approximately \$30,063, an increase of 2% over the 2021 rate. The ASC reimbursement rate for 2022 is approximately \$24,828, an increase of 2% over the 2021 rate.

Japan's Ministry of Health, Labour and Welfare ("MLHW") approved Inspire therapy to treat moderate to severe OSA in June 2018 and was formally added to the Japan National Health Insurance Payment Listing in June 2021. The first implants of Inspire therapy in Japan occurred in February 2022. In August 2020, the Australian Therapeutic Goods Administration approved Inspire therapy to treat moderate to severe OSA, and we are currently seeking reimbursement in Australia.

For the year ended December 31, 2021, 94.7% of our revenue was derived in the U.S. and 5.3% was derived outside of the U.S. No single customer accounted for more than 10% of our revenue.

Our marketing efforts during the first half of 2020 included refocused direct-to-consumer marketing strategies, which initially included a shift from radio and TV in our larger markets that were affected by COVID-19 towards more digital and TV in smaller markets. During the second quarter of 2020, and continuing through today, we resumed radio and TV initiatives in our larger markets as the impact of COVID-19 lessened in those areas. We continue to monitor the impacts of COVID-19 in each advertising market and may again change our advertising strategy on a market-by-market basis if needed. Further, our team has leveraged virtual tools, such as the new Inspire Sleep app released during the second quarter of 2020, to continue physician training and patient education. In January 2021, we began airing four new TV commercials and in January 2022, we purchased our first national television advertising spots and will begin airing more new TV commercials in the first quarter. We expect to continue to increase our digital and social media presence.

In early 2020, we started a call center concept, the Inspire Advisor Care Program ("ACP"). The primary purpose of this program is to assist patients with making a connection with a qualified healthcare provider based on their specific needs. As of December 31, 2021, approximately 550 of our U.S. centers were utilizing the ACP, up from approximately 180 centers at December 31, 2020. We plan to continue to increase the number of our U.S. centers using the ACP.

One of the many benefits of the ACP is the anecdotal feedback we are able to collect from patients during conversations with the ACP representatives. An example of this is the intelligence we have gathered that leads us to believe that we are beginning to see increased patient inquiries about Inspire as a result of the Philips Respironics CPAP recall. Following the recall announcement in July 2021, it took time for patients to learn about the recall, become educated about treatment alternatives and ultimately schedule an appointment with a healthcare provider to determine eligibility for an Inspire procedure. While we cannot quantify the impact from the recall, the feedback from ACP, as well as prior authorizations data and the Inspire app, all indicate increased patient flow as a result of the Philips recall. Looking ahead, we believe that there could be a sustained benefit to our business as a result of the recall.

We rely on third-party suppliers to manufacture our Inspire system and its components. Many of these suppliers are currently single source suppliers. We have experienced and continue to experience minor supply disruptions during the COVID pandemic, but have managed to avoid any significant supply and inventory issues. We seek to maintain higher levels of inventory to protect ourselves from supply interruptions, and, as a result, we are subject to the risk of inventory obsolescence and expiration, which could lead to inventory impairment charges. In the U.S., our products are shipped directly to our customers on a purchase order basis, primarily by a third-party vendor with a facility in Tennessee, although we do ship some products from our facility in Minnesota. Warehousing and shipping operations for our European customers are handled by a third-party vendor with a facility located in the Netherlands. Shipments of products to our Japanese distributor are handled from our facility in Minnesota. Customers do not have the right to return non-defective product, nor do we place product on consignment. Our sales representatives do not maintain trunk stock.

Since our inception in 2007, we have financed our operations primarily through sales of our Inspire system, private placements of our convertible preferred securities, amounts borrowed under our credit facility and equity offerings of our common stock. We have devoted significant resources to research and development activities related to our Inspire system, including clinical and regulatory initiatives to obtain marketing approval, and sales and marketing activities. For the year ended December 31, 2021, we generated revenue of \$233.4 million with a gross margin of 85.7% and a net loss of \$42.0 million, compared to revenue of \$115.4 million with a gross margin of 84.7% and a net loss of \$57.2 million for the year ended December 31, 2020, and revenue of \$82.1 million with

a gross margin of 83.4% and a net loss of \$33.2 million for the year ended December 31, 2019. Our accumulated deficit as of December 31, 2021 was \$279.4 million.

We have invested heavily in product development. Our research and development activities have been centered on driving continuous improvements to our Inspire therapy. We have also made significant investments in clinical studies to demonstrate the safety and efficacy of our Inspire therapy and to support regulatory submissions. We continue to make investments in research and development efforts to develop our next generation Inspire systems and support our future regulatory submissions for expanded indications and for new markets such as Europe, Japan, and Australia. For example, in December 2021, we received FDA approval for our Bluetooth-enabled patient remote control. In the first quarter of 2021, we received FDA approvals for both a new Inspire physician programmer platform and an improved two-incision surgical implant procedure that eliminates one incision with a revised placement of the pressure sensing lead. In May 2021, we received CE Mark approval in Europe for the two-incision implant procedure. In April 2020, we received FDA approval for an expanded age-range for Inspire therapy to include 18 to 21 year old patients. Japan's MLHW approved Inspire therapy to treat moderate to severe OSA in June 2018 and was formally added to the Japan National Health Insurance Payment Listing in June 2021. In August 2020, the Australian Therapeutic Goods Administration approved Inspire therapy to treat moderate to severe OSA.

We also continue to make significant investments building our sales and marketing organization by increasing the number of U.S. sales representatives and continuing our direct-to-consumer marketing efforts in existing and new markets throughout the U.S. and in Europe. During 2021, we activated 259 centers bringing the total to 684 U.S. medical centers implanting Inspire therapy as of December 31, 2021. Driven by the more favorable reimbursement environment, we have increased our focus on adding ASCs. At the end of 2021, ASCs made up 22% of our total U.S. implanting centers, up from 16% at the end of 2020. Additionally, we created 50 new sales territories during 2021, bringing the total to 157 U.S. territories as of December 31, 2021.

Because of these and other factors, we expect to continue to incur net losses for the next several years, and we may require substantial additional funding, which may include future equity and debt financings.

### **COVID-19 Pandemic Update**

Our business, operations and financial condition and results have been and may continue to be impacted by the COVID-19 pandemic. In 2020, we experienced significant reduction in revenue and product sales, as our customers were negatively impacted by the decline in the volume of elective procedures that resulted from the global healthcare system's response to COVID-19. As 2021 progressed, we observed a diminishing degree of COVID-related impacts to our reported revenue, although we believe there continues to be some adverse impact on our revenues. However, the extent to which the COVID-19 pandemic continues to impact our results of operations and financial condition will depend on future developments that are highly uncertain and cannot be predicted, including new information that may emerge concerning the severity and longevity of COVID-19 and its variants, the resurgence of COVID-19 in regions that have begun to recover from the initial impact of the pandemic, the impact of COVID-19 on economic activity, and the actions to contain its impact on public health and the global economy. For further information, refer to "Risk Factors" in Part II, Item 1A of this Annual Report on Form 10-K.

### **Components of Our Results of Operations**

#### **Revenue**

We derive primarily all of our revenue from the sale of our Inspire system to hospitals and ASCs in the U.S. and select countries in Europe. We recognize revenues from sales of our Inspire system when the customer obtains control of the product, which occurs at a point in time, either upon shipment of the product or receipt of the product, depending on shipment terms.

Our revenue has fluctuated, and may continue to fluctuate, from quarter to quarter due to a variety of factors. For example, we have historically experienced seasonality in our first and fourth quarters and have experienced adverse impacts on our revenue due to the COVID-19 pandemic.



### ***Cost of Goods Sold and Gross Margin***

Cost of goods sold consists primarily of acquisition costs for the components of the Inspire system, overhead costs, scrap, and inventory obsolescence, warranty replacement costs, as well as distribution-related expenses such as logistics and shipping costs, net of shipping costs charged to customers. The overhead costs include the cost of material procurement, depreciation expense for production equipment, and operations supervision and management personnel, including employee compensation, stock-based compensation, supplies, and travel. We expect cost of goods sold to increase or decrease in absolute dollars primarily as, and to the extent, our revenue grows or declines, respectively.

We calculate gross margin as gross profit divided by revenue. Our gross margin has been and we expect it will continue to be affected by a variety of factors, including manufacturing costs, the average selling price of our Inspire system, the implementation of cost-reduction strategies, inventory obsolescence costs, which generally occur when new generations of our Inspire system are introduced, and to a lesser extent the sales mix between the U.S. and Europe as our average selling price in the U.S. tends to be higher than in Europe. Our gross margin may increase slightly over the long term to the extent our production volumes increase and we receive discounts on the costs charged by our contract manufacturers, thereby reducing our per unit costs. However, our gross margin may fluctuate from quarter to quarter due to seasonality.

### ***Research and Development Expenses***

Research and development expenses consist primarily of product development, engineering, clinical studies to develop and support our products, regulatory expenses, quality assurance, testing, consulting services and other costs associated with the next generation versions of the Inspire system. These expenses include employee compensation, including stock-based compensation, supplies, materials, consulting, and travel expenses related to research and development programs. Additionally, these expenses include clinical trial management, payments to clinical investigators, data management and travel expenses for our various clinical trials.

We expect research and development expenses to increase in the future as we develop next generation versions of our Inspire system and continue to expand our clinical studies to further expand positive coverage policies from private commercial payors in the U.S. and enter into new markets including additional European countries and the Asia Pacific region. We expect research and development expenses as a percentage of revenue to vary over time depending on the level and timing of initiating new product development efforts and new clinical development activities.

### ***Selling, General and Administrative Expenses***

Selling, general and administrative ("SG&A") expenses consist primarily of compensation for personnel, including base salaries, stock-based compensation expense and commissions related to our sales organization, finance, information technology, and human resource functions, as well as spending related to marketing, sales operations, and training and reimbursement personnel. Other SG&A expenses include training physicians, travel expenses, advertising, direct-to-consumer promotional programs, conferences, trade shows and consulting services, professional services fees, audit fees, insurance costs and general corporate expenses, including facilities-related expenses.

We expect SG&A expenses to continue to increase as we expand our commercial infrastructure to both drive and support our planned growth in revenue and as we increase our headcount and expand administrative personnel to support our growth and operations as a public company including finance personnel and information technology services. Additionally, we anticipate an increase in our stock-based compensation expense with grants of restricted stock units, stock options, and shares of our common stock purchased pursuant to our employee stock purchase plan.

### ***Other Expense (Income), Net***

Other expense (income) consists primarily of interest expense payable under our credit facility and interest income.

## Results of Operations

### Year Ended December 31, 2021 Compared to Year Ended December 31, 2020

	Year Ended December 31,		Change	
	2021	2020	\$	%
(in thousands, except percentages)				
Revenue	\$ 233,394	\$115,381	\$ 118,013	102.3 %
Cost of goods sold	33,279	17,623	15,656	88.8 %
Gross profit	200,115	97,758	102,357	104.7 %
Gross margin	85.7 %	84.7 %		
Operating expenses:				
Research and development	37,350	26,092	11,258	43.1 %
Selling, general and administrative	202,615	127,874	74,741	58.4 %
Total operating expenses	239,965	153,966	85,999	55.9 %
Operating loss	(39,850)	(56,208)	16,358	(29.1)%
Other expense (income), net	2,120	880	1,240	140.9 %
Loss before income taxes	(41,970)	(57,088)	15,118	(26.5)%
Income taxes	72	115	(43)	(37.4)%
Net loss	\$ (42,042)	\$ (57,203)	\$ 15,161	(26.5)%

#### Revenue

Revenue increased \$118.0 million, or 102.3%, to \$233.4 million for the year ended December 31, 2021 compared to the year ended December 31, 2020. These results reflect an increase in sales of our Inspire system of \$114.9 million in the U.S. and an increase of \$3.1 million outside of the U.S., primarily in Germany. Revenue growth was primarily due to increased market penetration in existing territories, expansion into new territories, increased physician and patient awareness of our Inspire system, as well as the impact of the COVID-19 pandemic, which, as discussed below, hindered our performance during fiscal 2020.

Beginning in March 2020, our revenue growth in the U.S. and Europe was impacted by the COVID-19 pandemic, which disrupted our ability to access our clinician customers and their patients. Specifically, we saw healthcare facilities and clinics restricting access to their clinicians, reducing patient consultations and treatments, or closing temporarily due to COVID-19. As a result, beginning in the second week of March 2020, substantially all of our Inspire therapy procedures were postponed and numerous other cases, which had received prior authorization, were not able to be scheduled and, therefore were also postponed. Beginning in May 2020, surgical volumes began to increase steadily through the remainder of 2020, with most implanting centers performing procedures by October 2020. During the year ended December 31, 2021, resurgences of COVID-19 in various U.S. and European regions, as well as Japan, disrupted our ability to access our clinician customers and their patients, although to a lesser extent than in the same prior year period as surgical volumes generally returned to pre-pandemic levels by the end of the first quarter of 2021.

Revenue information by region is summarized as follows:

	Year Ended December 31,				Change	
	2021		2020			
	Amount	% of Revenue	Amount	% of Revenue	\$	%
(in thousands, except percentages)						
United States	\$ 220,976	94.7 %	\$ 106,108	92.0 %	\$ 114,868	108.3 %
Rest of world	12,418	5.3 %	9,273	8.0 %	3,145	33.9 %
Total revenue	\$ 233,394	100.0 %	\$ 115,381	100.0 %	\$ 118,013	102.3 %

Revenue generated in the U.S. was \$221.0 million for the year ended December 31, 2021, an increase of \$114.9 million, or 108.3%, over the year ended December 31, 2020. Revenue growth in the U.S. was primarily due to increased market penetration in existing territories, expansion into new territories, increased physician and patient awareness of our Inspire system, as well as the widespread shutdown of elective surgical procedures in 2020. As noted above, U.S. revenue during both periods was negatively impacted by the COVID-19 pandemic.

Revenue generated in outside of the U.S. was \$12.4 million in the year ended December 31, 2021, an increase of \$3.1 million, or 33.9%, over the year ended December 31, 2020. Revenue growth was primarily due to increased market penetration in existing territories, the expansion of our European sales representatives into new territories, our first sales to our Japanese distributor, and increased physician and patient awareness of our Inspire system during the year ended December 31, 2021, as well as the widespread shutdown of elective surgical procedures in 2020. The remainder of our increased revenue was due to favorable exchange rates. As noted above, European revenue during both periods was negatively impacted by the COVID-19 pandemic.

#### *Cost of Goods Sold and Gross Margin*

Cost of goods sold increased \$15.7 million, or 88.8%, to \$33.3 million for the year ended December 31, 2021 compared to \$17.6 million for the year ended December 31, 2020. The increase was primarily due to product costs associated with higher sales volume of our Inspire system.

Gross margin was 85.7% for the year ended December 31, 2021 compared to 84.7% for the year ended December 31, 2020. Gross margin for the year ended December 31, 2021 was higher primarily due to higher sales volumes and manufacturing efficiencies.

#### *Research and Development Expenses*

Research and development expenses increased \$11.3 million, or 43.1%, to \$37.4 million for the year ended December 31, 2021 compared to \$26.1 million for the year ended December 31, 2020. This change was primarily due to an increase of \$5.9 million of compensation and employee-related expenses, mainly as a result of increased headcount and stock-based compensation expense, \$5.9 million for ongoing research and development costs, including ongoing development of the Inspire Cloud and the next generation Inspire system, somewhat offset by a \$0.5 million decrease in regulatory submissions and clinical studies expenses.

#### *Selling, General and Administrative Expenses*

SG&A expenses increased \$74.7 million, or 58.4%, to \$202.6 million for the year ended December 31, 2021 compared to \$127.9 million for the year ended December 31, 2020. The primary driver of this change was an increase of \$46.4 million in compensation, including salaries, commissions, stock-based compensation, and other employee-related expenses, mainly as a result of increased headcount. In addition, marketing expenses increased \$24.6 million, primarily consisting of direct-to-consumer initiatives, including new TV advertisements, which began airing in January 2021 and the expansion of our Advisor Care Program call center. During the first part of 2020, we initially refocused our direct-to-consumer marketing strategies by shifting from radio and TV in our larger markets that were affected by COVID-19 towards more digital and TV in smaller markets. During the second quarter of 2020, we resumed radio and TV initiatives in our larger markets as the impact of COVID-19

lessened in those areas. Further, our team leveraged virtual tools, including the new Inspire Sleep app released in the second quarter of 2020, and telemedicine to continue physician training and patient education. Other drivers of the change to SG&A expenses included an increase of \$3.7 million primarily due to office rent, insurance costs, consulting fees and banking fees.

#### *Other Expense (Income), Net*

Other expense (income), net changed by \$1.2 million, or 140.9%, to \$2.1 million of expense for the year ended December 31, 2021 compared to \$0.9 million of expense for the year ended December 31, 2020. This change was primarily due to a decrease of \$1.0 million in interest income due to lower interest rates on our cash, cash equivalents and investments balances and a \$0.3 million decrease in gains on investments and foreign currency translation.

#### *Income Taxes*

We recorded a \$0.1 million provision for income taxes for the both of the years ended December 31, 2021 and 2020.

#### ***Year Ended December 31, 2020 Compared to Year Ended December 31, 2019***

For a discussion of our results of operations for the year ended December 31, 2020, including a year-to-year comparison between 2020 and 2019, refer to Part II, Item 7, "Management's Discussion and Analysis of Financial Condition and Results of Operations" in our Annual Report Form 10-K for the year ended December 31, 2020.

#### **Liquidity and Capital Resources**

As of December 31, 2021, we had cash, cash equivalents and investments of \$224.4 million, a decrease of \$10.0 million from \$234.4 million as of December 31, 2020. Working capital totaled \$227.2 million as of December 31, 2021, a decrease of \$21.9 million from December 31, 2020. We define working capital as current assets less current liabilities. The decrease in working capital was primarily due to a \$19.9 million decrease in cash, cash equivalents and short term investments, \$9.9 million of which was used to support operations, and \$10.0 million of which was moved into long term investments. Working capital also decreased with the movement of \$9.2 million of our long-term debt into current liabilities, as the first principal payments become due in April 2022, as well as a \$6.9 million increase in accrued expenses primarily due to accrued compensation, and a \$4.5 million increase in accounts payable, generally due to our increased business volume year-over-year, increased headcount from the prior year and the costs to support the growth of our operations. The decrease in working capital was partially offset by an increase of \$9.1 million in accounts receivable due to higher sales, an increase of \$8.8 million in inventory balances which increased to meet increased sales and to establish safety stock to avoid inventory shortages in the event of COVID-related production or supply issues, and a \$0.7 million increase in prepaid expense and other current assets.

We proactively manage our access to capital to support liquidity and continued growth. Our sources of capital include sales of our Inspire system, borrowings under credit facilities and registered offerings of our common stock. At December 31, 2021, we had \$24.5 million of outstanding borrowings under our credit facility and no borrowings remain available under this credit facility. We will begin paying principal payments on the borrowings in April 2022, and the scheduled maturity date of the facility is March 2024. We were in compliance with all covenants under the credit facility as of December 31, 2021. See Note 5 to our audited financial statements for additional information on our credit facility.

The primary objective of our investment activities is to preserve our capital for the purpose of funding operations while at the same time maximizing the income we receive from our investments without significantly increasing risk or decreasing availability. To achieve these objectives, our investment policy allows us to maintain a portfolio of certain types of debt securities issued by the U.S. government and its agencies, corporations with investment-grade credit ratings, or commercial paper and money market funds issued by the highest quality financial and non-financial companies. At December 31, 2021, we had \$9.9 million of investments in U.S. government securities, and no investments with a contractual maturity over two years.

In 2021, our SG&A expenditures increased significantly over 2020 levels, and we anticipate further increases in 2022. Our SG&A expenditures, primarily for increasing headcount and advertising, may exceed any associated increases in revenues, and therefore would reduce our cash flow from operations. We also anticipate R&D expenses will continue to be significant in fiscal 2022, primarily related to the ongoing development of the Inspire Cloud and the next generation Inspire system.

We spent \$4.7 million on purchases of property and equipment in 2021, mainly on testing systems, production equipment, and tooling. We anticipate further capital expenditures in 2022, primarily for additional equipment and tooling.

We believe that our existing cash and cash equivalents and investments, which totaled \$224.4 million as of December 31, 2021, together with cash flow from operations, will provide liquidity sufficient to meet our cash needs and fund our operations and planned capital expenditures for at least the next 12 months. There can be no assurance, however, that our business will continue to generate cash flows at historic levels.

Beyond fiscal 2022, our cash requirements will depend extensively on the timing of market introduction, and extent of market acceptance of, our Inspire system. Our long-term cash requirements also will be significantly impacted by the level of our investment in commercialization, entry into new markets such as Australia and Singapore, and whether we make strategic acquisitions. We cannot accurately predict our long-term cash requirements at this time. Additionally, The COVID-19 pandemic has negatively impacted the global economy, disrupted global supply chains and created significant volatility and disruption of financial markets. An extended period of global supply chain and economic disruption could materially affect our business, results of operations, access to sources of liquidity and financial condition. We may seek additional sources of liquidity and capital resources through additional securities offerings or through borrowings under a new credit facility. There can be no assurance that such transactions will be available to us on favorable terms, if at all.

Below is a summary of short-term and long-term anticipated cash requirements under contractual obligations existing as of December 31, 2021.

(\$ in thousands)	As of December 31, 2021		
	Total	Fiscal 2022	After Fiscal 2022
Recorded contractual obligations:			
Long-term debt principal payments <sup>(1)</sup>	\$ 24,500	\$ 9,188	\$ 15,312
Operating leases <sup>(2)</sup>	10,999	725	10,274
Unrecorded contractual obligations:			
Long-term debt interest and final payment fee payments <sup>(3)</sup>	3,295	1,651	1,644
Purchase obligations <sup>(4)</sup>	114,632	70,427	44,205
<b>Total</b>	<b>\$ 153,426</b>	<b>\$ 81,991</b>	<b>\$ 71,435</b>

<sup>(1)</sup> Represents principal payments only. See Note 5 to our audited financial statements for additional information.

<sup>(2)</sup> See Note 4 to our audited financial statements.

<sup>(3)</sup> Variable interest is assumed at December 31, 2021 rates. Under the terms of the loan and security agreement, a final payment fee of 3.50% is due at the earlier of maturity or prepayment. This amount is included in the table above assuming the final payment is made at debt maturity in March 2024. See Note 5 to our audited financial statements for additional information.

<sup>(4)</sup> Primarily purchase obligations to suppliers for inventory.

As of December 31, 2021, we did not have any off-balance sheet arrangements that have, or are reasonably likely to have, a current or future material effect on our financial condition, results of operations, liquidity, capital expenditures or capital resources.

## Cash Flows

The following table presents a summary of our cash flow for the periods indicated:

	Year Ended December 31,	
	2021	2020
	(in thousands)	
Net cash provided by (used in):		
Operating activities	\$ (20,119)	\$ (53,045)
Investing activities	29,139	86,627
Financing activities	14,948	134,065
Effect of exchange rate on cash	(19)	11
Increase in cash and cash equivalents	<u>\$ 23,949</u>	<u>\$ 167,658</u>

### Operating Activities

Net cash used in operating activities was \$20.1 million for 2021 and consisted of a net loss of \$42.0 million, non-cash charges of \$28.8 million, and increase in net operating assets of \$6.9 million. The non-cash charges consisted primarily of stock-based compensation, which increased mainly as a result of granting more stock options to employees at a higher fair market value. The remainder of the non-cash charges included depreciation and amortization, non-cash lease expense, stock issued for services rendered, accretion of the debt discount, and amortization of the investment premium, and other, net. Operating assets includes inventories, which increased due to manufacturing of systems inventory to meet increased sales and to establish safety stock to avoid inventory shortages in the event of COVID-related production or supply issues. Operating assets also includes accounts receivable which increased due to higher sales, and prepaid expenses and other current assets which increased primarily due to prepaid insurance. Operating liabilities includes accrued expenses, which increased primarily due to accrued compensation, and accounts payable, which increased generally due to our increased business volume year-over-year and the costs to support the growth of our operations.

Net cash used in operating activities was \$53.0 million for 2020 and consisted of a net loss of \$57.2 million, an increase in net operating assets of \$10.9 million, and non-cash charges of \$15.0 million. The non-cash charges consisted of stock-based compensation, non-cash lease expense, depreciation and amortization, stock issued for services rendered, and accretion of the debt discount, offset by the non-cash income related to the accretion of the investment discount, and other, net. Operating assets includes accounts receivable, which increased due to higher sales, and inventories, which increased due to continued manufacturing of systems inventory as sales volumes increase. Operating liabilities includes accounts payable which increased generally due to the costs to support the growth of our operations, including compensation and personnel-related costs. Accrued expenses also increased due to the accrued compensation for annual bonuses to be paid during the first quarter of 2021.

### Investing Activities

Net cash provided by investing activities for 2021 was \$29.1 million and consisted primarily of proceeds from sales or maturities of investments of \$43.8 million, partially offset by purchases of investments of \$10.0 million. Purchases of property and equipment of \$4.7 million, mainly for manufacturing test systems, production equipment, and tooling, comprised the remainder of the investing activities.

Net cash provided by investing activities for 2020 was \$86.6 million and consisted primarily of proceeds from sales or maturities of investments of \$141.8 million, partially offset by purchases of investments of \$52.7 million and purchases of property and equipment of \$2.5 million.

### Financing Activities

Net cash provided by financing activities was \$14.9 million for 2021 and consisted of \$11.5 million in proceeds from the exercise of stock options and \$3.5 million in proceeds from the issuance of common stock from our employee stock purchase plan ("ESPP").

Net cash provided by financing activities was \$134.1 million for 2020 and consisted primarily of \$124.7 million in proceeds from our follow-on offering in April 2020. The remainder of the cash provided by financing activities resulted from \$7.1 million in proceeds from the exercise of stock options and \$2.4 million in proceeds from the issuance of common stock from our ESPP.

### **Critical Accounting Estimates**

The preparation of the financial statements in conformity with accounting principles generally accepted in the U.S. requires management to make estimates and assumptions that affect the amounts reported in the audited financial statements and accompanying notes included elsewhere in this Annual Report on Form 10-K. We believe that such estimates have been based on reasonable and supportable assumptions and the resulting estimates are reasonable for use in the preparation of the audited financial statements. Actual results could differ from these estimates.

The following area requires management estimates, assumptions, and judgments:

#### ***Inventories***

Inventories are valued at the lower of cost or net realizable value, computed on a first-in, first out basis. We estimate the recoverability of our inventory by reference to internal estimates of future demands and product life cycles, including expiration of inventory prior to sale. We regularly review inventory quantities on-hand for excess and obsolete inventory and, when circumstances indicate, incur charges to write down inventories to their net realizable value. The determination of a reserve for excess and obsolete inventory involves management exercising judgment to determine the required reserve, considering future demand, product life cycles, introduction of new products and current market conditions. Although we make every effort to ensure the accuracy of our forecasts of future product demand, any significant unanticipated changes in demand could have a significant impact on the carrying value of our inventories and reported operating results. The net inventory balance was \$17.2 million and \$8.5 million as of December 31, 2021 and 2020, respectively. The reserve for excess and obsolete inventory was \$0.3 million and \$0.1 million as of December 31, 2021 and 2020, respectively.

### **Recent Accounting Pronouncements**

A discussion of recent accounting pronouncements is included in Note 2 to our financial statements contained in this Annual Report on Form 10-K.

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

### ***Interest Rate Risk***

The risk associated with fluctuating interest rates is primarily limited to our cash equivalents which are carried at quoted market prices and our short-term investments. We do not currently use or plan to use financial derivatives in our investment portfolio. The interest rate for our outstanding debt is variable. A hypothetical 1% change in interest rates during any of the periods presented would not have had a material impact on our consolidated financial statements.

### ***Credit Risk***

As of December 31, 2021 and 2020, our cash, cash equivalents, and investments were maintained with two financial institutions in the U.S. We believe these institutions have sufficient assets and liquidity to conduct their operations in the ordinary course of business with little or no credit risk to us, however our cash balances were in excess of insured limits.

The primary objective of our investment activities is to preserve our capital for the purpose of funding operations while at the same time maximizing the income we receive from our investments without significantly increasing risk or decreasing availability. To achieve these objectives, our investment policy allows us to maintain a portfolio

of certain types of debt securities issued by the U.S. government and its agencies, corporations with investment-grade credit ratings, or commercial paper and money market funds issued by the highest quality financial and non-financial companies. We place restrictions on maturities and concentration by type and issuer. We are exposed to credit risk in the event of a default by the issuers of these securities to the extent recorded on the balance sheets. See Note 2 to our financial statements contained in this Annual Report on Form 10-K for additional information on our cash equivalents and available-for-sale marketable securities.

Our accounts receivable primarily relate to revenue from the sale of our Inspire system to hospitals in the U.S. and Europe, primarily in Germany. No single customer represented more than 10% of our accounts receivable as of December 31, 2021 or 2020.

### ***Foreign Currency Risk***

The majority of our business is currently conducted in U.S. dollars. Any transactions that may be conducted in foreign currencies are not expected to have a material effect on our results of operations, financial position or cash flows.

### ***Inflation Risk***

Inflationary factors, such as increases in our cost of goods sold and selling and operating expenses, may adversely affect our operating results. Although we do not believe that inflation has had a material impact on our financial position or results of operations to date, a high rate of inflation in the future may have an adverse effect on our ability to maintain and increase our gross margin and selling and marketing and operating expenses as a percentage of our revenue if the selling prices of our products do not increase as much as or more than these increased costs.



## **Item 8. Financial Statements and Supplementary Data.**

### **Report of Independent Registered Public Accounting Firm**

To the Stockholders and the Board of Directors of Inspire Medical Systems, Inc.

#### **Opinion on the Financial Statements**

We have audited the accompanying balance sheets of Inspire Medical Systems, Inc. (the Company) as of December 31, 2021 and 2020, the related statements of operations and comprehensive loss, stockholders' equity and cash flows for each of the three years in the period ended December 31, 2021, and the related notes (collectively referred to as the "financial statements"). In our opinion, the financial statements present fairly, in all material respects, the financial position of the Company at December 31, 2021 and 2020, and the results of its operations and its cash flows for each of the three years in the period ended December 31, 2021, in conformity with U.S. generally accepted accounting principles.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States) (PCAOB), the Company's internal control over financial reporting as of December 31, 2021, based on criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework), and our report dated February 15, 2022 expressed an unqualified opinion thereon.

#### **Basis for Opinion**

These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on the Company's financial statements based on our audits. We are a public accounting firm registered with the PCAOB and are required to be independent with respect to the Company in accordance with the U.S. federal securities laws and the applicable rules and regulations of the Securities and Exchange Commission and the PCAOB.

We conducted our audits in accordance with the standards of the PCAOB. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement, whether due to error or fraud. Our audits included performing procedures to assess the risks of material misstatement of the financial statements, whether due to error or fraud, and performing procedures that respond to those risks. Such procedures included examining, on a test basis, evidence regarding the amounts and disclosures in the financial statements. Our audits also included evaluating the accounting principles used and significant estimates made by management, as well as evaluating the overall presentation of the financial statements. We believe that our audits provide a reasonable basis for our opinion.

#### **Critical Audit Matter**

The critical audit matter communicated below is a matter arising from the current period audit of the financial statements that was communicated or required to be communicated to the audit committee and that: (1) relates to accounts or disclosures that are material to the financial statements and (2) involved our especially challenging, subjective or complex judgments. The communication of the critical audit matter does not alter in any way our opinion on the financial statements, taken as a whole, and we are not, by communicating the critical audit matter

below, providing a separate opinion on the critical audit matter or on the account or disclosure to which it relates.

***Inventory valuation reserves***

***Description of the Matter***

At December 31, 2021, the Company's net inventory balance was \$17.2 million. As explained in Note 2 to the financial statements, the determination of a reserve for excess and obsolete inventory involves management exercising judgment to determine the required reserve, considering future demand, product life cycles, introduction of new products and current market conditions.

Auditing management's estimate for excess and obsolete inventory involved subjective auditor judgment because of the assumptions and judgments used to calculate the inventory valuation reserve, including consideration of the timing of the introduction of new products and current market conditions. In particular, the excess and obsolete inventory calculations are sensitive to significant assumptions, including forecasted customer demand, technological and/or market obsolescence, and possible alternative uses.

***How We Addressed the Matter in Our Audit***

We obtained an understanding, evaluated the design, and tested the operating effectiveness of controls over the Company's process to determine the amount of the Company's reserve for excess and obsolete inventory. This included controls over the Company's review of the significant assumptions underlying the reserve estimate.

To test the adequacy of the Company's inventory valuation reserve, we performed audit procedures that included, among others, testing the accuracy and completeness of the underlying data used in the estimation calculations and evaluating significant assumptions, specifically forecasted customer demand, technological and/or market obsolescence, and possible alternative uses. We evaluated management's ability to accurately estimate the amount of excess and obsolete inventory by comparing actual inventory write-off activity in recent years to management's prior year estimates of the inventory valuation reserve. We also audited management's calculation of the inventory valuation reserve by testing the mathematical accuracy of the Company's reserve calculation.

/s/Ernst & Young LLP

We have served as the Company's auditor since 2015.

Minneapolis, Minnesota  
February 15, 2022

**INSPIRE MEDICAL SYSTEMS, INC.**

**BALANCE SHEETS**

(in thousands, except share and per share amounts)

	<b>December 31,</b>	
	<b>2021</b>	<b>2020</b>
<b>Assets</b>		
Current assets:		
Cash and cash equivalents	\$ 214,467	\$ 190,518
Investments, short-term	—	43,844
Accounts receivable, net of allowance for credit losses of \$99 and \$42, respectively	34,179	25,063
Inventories	17,231	8,479
Prepaid expenses and other current assets	2,660	1,965
Total current assets	268,537	269,869
Investments, long-term	9,938	—
Property and equipment, net	8,486	5,311
Operating lease right-of-use asset	7,919	5,805
Other non-current assets	204	204
Total assets	<u>\$ 295,084</u>	<u>\$ 281,189</u>
<b>Liabilities and stockholders' equity</b>		
Current liabilities:		
Accounts payable	\$ 11,665	\$ 7,209
Accrued expenses	20,454	13,516
Notes payable, current portion	9,188	—
Total current liabilities	41,307	20,725
Notes payable, non-current portion	15,799	24,746
Operating lease liability, non-current portion	8,796	5,886
Other non-current liability	134	85
Total liabilities	66,036	51,442
Stockholders' equity		
Preferred Stock, \$0.001 par value, 10,000,000 shares authorized; no shares issued and outstanding	—	—
Common Stock, \$0.001 par value, 200,000,000 shares authorized; 27,416,106 and 27,069,276 shares issued and outstanding at December 31, 2021 and 2020, respectively	27	27
Additional paid-in capital	508,465	467,038
Accumulated other comprehensive (loss) income	(55)	29
Accumulated deficit	(279,389)	(237,347)
Total stockholders' equity	229,048	229,747
Total liabilities and stockholders' equity	<u>\$ 295,084</u>	<u>\$ 281,189</u>

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

**INSPIRE MEDICAL SYSTEMS, INC.**  
**STATEMENTS OF OPERATIONS AND COMPREHENSIVE LOSS**  
(in thousands, except share and per share amounts)

	Year Ended December 31,		
	2021	2020	2019
Revenue	\$ 233,394	\$ 115,381	\$ 82,050
Cost of goods sold	33,279	17,623	13,643
Gross profit	200,115	97,758	68,407
Operating expenses:			
Research and development	37,350	26,092	12,839
Selling, general and administrative	202,615	127,874	90,465
Total operating expenses	239,965	153,966	103,304
Operating loss	(39,850)	(56,208)	(34,897)
Other expense (income):			
Interest income	(125)	(1,092)	(3,801)
Interest expense	2,128	2,117	2,119
Other expense (income), net	117	(145)	(12)
Total other expense (income)	2,120	880	(1,694)
Loss before income taxes	(41,970)	(57,088)	(33,203)
Income taxes	72	115	40
Net loss	(42,042)	(57,203)	(33,243)
Other comprehensive loss:			
Unrealized (loss) gain on investments	(84)	(73)	154
Total comprehensive loss	<u>\$ (42,126)</u>	<u>\$ (57,276)</u>	<u>\$ (33,089)</u>
Net loss per share, basic and diluted	<u>\$ (1.54)</u>	<u>\$ (2.19)</u>	<u>\$ (1.40)</u>
Weighted average common shares used to compute net loss per share, basic and diluted	27,262,979	26,073,418	23,804,452

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

**INSPIRE MEDICAL SYSTEMS, INC.**  
**STATEMENTS OF STOCKHOLDERS' EQUITY**  
(in thousands, except share amounts)

	<u>Common Stock</u>		<u>Additional Paid-in Capital</u>	<u>Accumulated Other Comprehensive (Loss) Income</u>	<u>Accumulated Deficit</u>	<u>Total Stockholders' Equity</u>
	<u>Shares</u>	<u>Amount</u>				
Balance at December 31, 2018	23,401,675	\$ 23	\$ 310,941	\$ (52)	\$ (146,913)	\$ 163,999
Stock options and warrants exercised	669,719	1	1,139	—	—	1,140
Issuance of common stock	4,206	—	233	—	—	233
Issuance of common stock for employee stock purchase plan	31,750	—	1,355	—	—	1,355
Stock-based compensation expense	—	—	6,197	—	—	6,197
Other comprehensive income	—	—	—	154	—	154
Net loss	—	—	—	—	(33,243)	(33,243)
Balance at December 31, 2019	24,107,350	24	319,865	102	(180,156)	139,835
Stock options exercised	624,315	—	7,050	—	—	7,050
Issuance of common stock	3,378	—	294	—	—	294
Sale of common stock from follow-on public offering, net of offering expenses	2,300,000	3	124,651	—	—	124,654
Issuance of common stock for employee stock purchase plan	34,233	—	2,361	—	—	2,361
Stock-based compensation expense	—	—	12,817	—	—	12,817
Other comprehensive loss	—	—	—	(73)	—	(73)
Adoption of ASU 2016-13, Financial Instruments - Credit Losses	—	—	—	—	12	12
Net loss	—	—	—	—	(57,203)	(57,203)
Balance at December 31, 2020	27,069,276	27	467,038	29	(237,347)	229,747
Stock options exercised	323,860	—	11,476	—	—	11,476
Issuance of common stock	1,463	—	301	—	—	301
Issuance of common stock for employee stock purchase plan	21,507	—	3,472	—	—	3,472
Stock-based compensation expense	—	—	26,178	—	—	26,178
Other comprehensive loss	—	—	—	(84)	—	(84)
Net loss	—	—	—	—	(42,042)	(42,042)
Balance at December 31, 2021	<u>27,416,106</u>	<u>\$ 27</u>	<u>\$ 508,465</u>	<u>\$ (55)</u>	<u>\$ (279,389)</u>	<u>\$ 229,048</u>

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

**INSPIRE MEDICAL SYSTEMS, INC.**  
**STATEMENTS OF CASH FLOWS**  
(in thousands)

	Year Ended December 31,		
	2021	2020	2019
<b>Operating activities</b>			
Net loss	\$ (42,042)	\$ (57,203)	\$ (33,243)
Adjustments to reconcile net loss:			
Depreciation and amortization	1,218	840	495
Amortization (accretion) of investment premium (discount)	14	(22)	(983)
Accretion of debt discount	240	224	268
Non-cash lease expense	771	972	904
Stock-based compensation expense	26,178	12,817	6,197
Non-cash stock issuance for services rendered	301	294	233
Other, net	56	(89)	(143)
Changes in operating assets and liabilities:			
Accounts receivable	(9,244)	(11,842)	(6,480)
Inventories	(8,752)	(2,645)	(3,167)
Prepaid expenses and other assets	(696)	416	(905)
Accounts payable	4,779	2,048	1,034
Accrued expenses and other liabilities	7,058	1,145	2,944
Net cash used in operating activities	(20,119)	(53,045)	(32,846)
<b>Investing activities</b>			
Purchases of property and equipment	(4,668)	(2,455)	(2,739)
Purchases of investments	(9,993)	(52,721)	(178,074)
Proceeds from sales or maturities of investments	43,800	141,803	137,254
Net cash provided by (used in) investing activities	29,139	86,627	(43,559)
<b>Financing activities</b>			
Payment of debt fees	—	—	(531)
Proceeds from the exercise of stock options and warrants	11,476	7,050	1,140
Proceeds from sale of common stock	—	124,654	—
Proceeds from the issuance of common stock from employee stock purchase plan	3,472	2,361	1,355
Net cash provided by financing activities	14,948	134,065	1,964
Effect of exchange rate on cash	(19)	11	13
Increase (decrease) in cash and cash equivalents	23,949	167,658	(74,428)
Cash and cash equivalents at beginning of year	190,518	22,860	97,288
Cash and cash equivalents at end of year	<u>\$ 214,467</u>	<u>\$ 190,518</u>	<u>\$ 22,860</u>
<b>Supplemental cash flow information</b>			
Cash paid for interest	\$ 1,888	\$ 1,893	\$ 2,033
Change in property and equipment acquired but not yet paid	274	816	—

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

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
(Table amounts in thousands, except share and per share amounts)

## **1. Organization**

### **Description of Business**

Inspire Medical Systems, Inc. is a medical technology company focused on the development and commercialization of innovative, minimally invasive solutions for patients with obstructive sleep apnea ("OSA"). Our proprietary Inspire system is the first and only United States ("U.S.") Food and Drug Administration ("FDA") approved neurostimulation technology that provides a safe and effective treatment for moderate to severe OSA. Inspire therapy received premarket approval ("PMA") from the FDA in April 2014 and has been commercially available in certain European markets since November 2011. Japan's Ministry of Health, Labour and Welfare ("MLHW") approved Inspire therapy to treat moderate to severe OSA in June 2018 and was formally added to the Japan National Health Insurance Payment Listing in June 2021. In August 2020, the Australian Therapeutic Goods Administration approved Inspire therapy to treat moderate to severe OSA, and we are currently seeking reimbursement coverage in Australia.

## **2. Summary of Significant Accounting Policies**

### **Basis of Presentation**

The financial statements have been prepared in accordance with U.S. generally accepted accounting principles ("U.S. GAAP") and pursuant to the rules and regulations of the Securities and Exchange Commission ("SEC").

In the opinion of management, all adjustments, consisting of only normal recurring adjustments that are necessary to present fairly the financial position, results of operations, and cash flows have been made. The results of operations for the year ended December 31, 2021 are not necessarily indicative of the operating results for any future periods.

### **Use of Estimates**

The preparation of financial statements in conformity with U.S. GAAP requires management to make estimates and assumptions that affect the amounts and disclosures reported in the financial statements. We use significant judgment when making estimates related to the allowance for credit losses, inventory reserves, warranty reserves, and stock-based awards. We base our estimates on historical experience and on various other assumptions that are believed to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Actual results could differ from those estimates.

### **Cash and Cash Equivalents**

We consider all highly liquid securities, readily convertible to cash, that mature within 90 days or less from the date of purchase to be cash equivalents. The carrying amount reported in the balance sheets for cash is cost, which approximates fair value.

### **Foreign Currency**

Sales and expenses denominated in foreign currencies are translated at average exchange rates in effect throughout the year. Foreign currency transaction gains and losses are included in other expense (income), net in the statements of operations and comprehensive loss. Assets and liabilities of foreign operations are remeasured at period-end exchange rates with the impacts of foreign currency remeasurement recognized in other expense (income), net in the statements of operations and comprehensive loss.

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
**(Table amounts in thousands, except share and per share amounts)**

**Investments**

At December 31, 2021, our long-term investments consisted of U.S. government securities. At December 31, 2020, our short-term investments consisted of commercial paper, corporate bonds, and U.S. government securities. Investments are reported at their estimated fair market values which are based on quoted, active or inactive market prices when available. Any unrealized gains and losses due to interest rate fluctuations and other external factors are reported as a separate component of accumulated other comprehensive (loss) income. We had \$0.1 million of unrecognized loss and approximately \$0 of unrecognized income in our accumulated other comprehensive (loss) income balance at December 31, 2021 and 2020, respectively. Any realized gains and losses are calculated on the specific identification method and reported net in other expense (income) in the statements of operations and comprehensive loss. For the years ended December 31, 2021 and 2020, we recognized \$0 and \$0.1 million of gains, net, respectively.

We recognize expected credit losses on investments in accordance with Accounting Standards Update ("ASU"), ASU 2016-13, *Financial Instruments - Credit Losses* ("ASU 2016-13"), which we adopted effective January 1, 2020 using the modified retrospective approach. The adoption of ASU 2016-13 did not have a material impact on the amount and timing of credit losses recognized in our financial statements.

We reassess our estimated credit losses on investments each reporting period. U.S. government securities and cash equivalents are under a "zero-loss exception" for credit losses, meaning no credit loss risk calculation is necessary on those instruments due to the exceptionally low rate of default, which continues to decrease as the securities approach maturity, which for us is no longer than two years. For non-U.S. government securities, we use a discounted cash flow approach to calculate expected credit losses using estimated default rates based upon historical loss data, current conditions, as well as expectations of future economic conditions. We record changes in the allowance for credit losses for available-for-sale debt securities with a corresponding adjustment in credit loss expense on the statement of operations and comprehensive loss. No reversal of a previously recorded allowance for credit losses may be made to an amount below zero. The total allowance for credit losses was \$0 at both December 31, 2021 and 2020.

**Fair Value of Financial Instruments**

We measure certain financial assets and liabilities at fair value on a recurring basis, including cash equivalents and investments. Fair value is an exit price, representing the amount that would be received to sell an asset or paid to transfer a liability in an orderly transaction between market participants. As such, fair value is a market-based measurement that should be determined based on assumptions that market participants would use in pricing an asset or a liability. A three-tier fair value hierarchy is established as a basis for considering such assumptions and for inputs used in the valuation methodologies in measuring fair value:

Level 1 — Observable inputs, such as quoted prices (unadjusted) for identical assets or liabilities in active markets.

Level 2 — Inputs other than quoted prices in active markets for identical assets and liabilities that are observable either directly or indirectly for substantially the full term of the asset or liability.

Level 3 — Unobservable inputs that are supported by little or no market activities, which would require us to develop our own assumptions.

We use the methods and assumptions described below in determining the fair value of our financial instruments.

*Money market funds:* Fair values of money market funds are based on quoted market prices in active markets. These are included as Level 1 measurements in the tables below.

*Commercial paper:* Short-term, highly liquid investments are included as a Level 2 measurement in the table below.



**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
(Table amounts in thousands, except share and per share amounts)

*Corporate bonds:* Consists of short- and long-term notes and bonds with various yields. These are included as a Level 2 measurement in the table below.

*U.S. government securities:* Consists of U.S. government Treasury bills with original maturities of less than two years. These are included as a Level 1 measurement in the tables below.

The following tables sets forth by level within the fair value hierarchy our assets that are measured on a recurring basis and reported at fair value as of December 31, 2021 and 2020. Assets are classified in their entirety based on the lowest level of input that is significant to the fair value measurement.

Fair Value Measurements as of December 31, 2021				
	Estimated Fair Value	Level 1	Level 2	Level 3
Cash equivalents:				
Money market funds	\$ 189,369	\$ 189,369	\$ —	\$ —
Total cash equivalents	189,369	189,369	—	—
Investments:				
U.S. government securities	9,938	9,938	—	—
Total investments	9,938	9,938	—	—
Total cash equivalents and investments	\$ 199,307	\$ 199,307	\$ —	\$ —
Fair Value Measurements as of December 31, 2020				
	Estimated Fair Value	Level 1	Level 2	Level 3
Cash equivalents:				
Money market funds	\$ 179,389	\$ 179,389	\$ —	\$ —
Total cash equivalents	179,389	179,389	—	—
Investments:				
Commercial paper	13,275	—	13,275	—
Corporate bonds	6,540	—	6,540	—
U.S. government securities	24,029	24,029	—	—
Total investments	43,844	24,029	19,815	—
Total cash equivalents and investments	\$ 223,233	\$ 203,418	\$ 19,815	\$ —

There were no transfers between levels during the years ended December 31, 2021 and 2020.

### Concentration of Credit Risk

Financial instruments, which potentially subject us to concentrations of credit risk, consist principally of cash equivalents, investments, and accounts receivable.

Our investment policy limits investments to certain types of debt securities issued by the U.S. government and its agencies, corporations with investment-grade credit ratings, or commercial paper and money market funds issued by the highest quality financial and non-financial companies. We place restrictions on maturities and concentration by type and issuer. We are exposed to credit risk in the event of a default by the issuers of these securities to the extent recorded on the balance sheets. However, as of December 31, 2021 and 2020, we limited our credit risk associated with cash equivalents by placing investments with banks we believe are highly creditworthy.

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
**(Table amounts in thousands, except share and per share amounts)**

We believe that the credit risk in our accounts receivable is mitigated by our credit evaluation process, relatively short collection terms, and dispersion of our customer base. We generally do not require collateral, and losses on accounts receivable have historically not been significant.

**Accounts Receivable and Allowance for Expected Credit Losses**

Trade accounts receivable are recorded at the invoiced amount and do not bear interest. Customer credit terms are established prior to shipment with the general standard being net 30 days. Collateral or any other security to support payment of these receivables generally is not required.

We recognize expected credit losses on accounts receivable in accordance with ASU 2016-13, *Financial Instruments - Credit Losses* ("ASU 2016-13"), which we adopted effective January 1, 2020 using the modified retrospective approach through a cumulative-effect adjustment to retained earnings. The adoption of ASU 2016-13 did not have a material impact on the amount and timing of credit losses recognized in our financial statements.

Each reporting period, we estimate the credit loss related to accounts receivable based on a migration analysis of accounts grouped by individual receivables delinquency status, and apply our historic loss rate adjusted for management's assumption of future market conditions. Any change in the allowance from new receivables acquired, or changes due to credit deterioration on previously existing receivables, is recorded in selling, general and administrative expenses. Write-offs of receivables considered uncollectible are deducted from the allowance. Specific accounts receivable are written-off once a determination is made that the amount is uncollectible. The write-off is recorded in the period in which the account receivable is deemed uncollectible. Recoveries are recognized when received and as a direct credit to earnings or as a reduction to the allowance for credit losses (which would indirectly reduce the loss by decreasing bad debt expense).

**Inventories**

Inventories are valued at the lower of cost or net realizable value, computed on a first-in, first-out basis, and consisted of the following:

	December 31,	
	2021	2020
Raw materials	\$ 3,119	\$ 892
Finished goods	14,112	7,587
Total inventories, net of reserves	<u>\$ 17,231</u>	<u>\$ 8,479</u>

We regularly review inventory quantities on-hand for excess and obsolete inventory and, when circumstances indicate, incur charges to write down inventories to their net realizable value. The determination of a reserve for excess and obsolete inventory involves management exercising judgment to determine the required reserve, considering future demand, product life cycles, introduction of new products and current market conditions. The reserve for excess and obsolete inventory was \$0.3 million and \$0.1 million as of December 31, 2021 and 2020, respectively.

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
(Table amounts in thousands, except share and per share amounts)

**Property and Equipment**

Property and equipment are stated at cost, less accumulated depreciation and amortization and consisted of the following:

	<b>December 31,</b>	
	<b>2021</b>	<b>2020</b>
Computer equipment and software	\$ 1,397	\$ 1,305
Manufacturing equipment	4,436	2,285
Other equipment	249	249
Leasehold improvements	281	192
Construction in process	5,175	3,125
Property and equipment, cost	11,538	7,156
Less: accumulated depreciation and amortization	(3,052)	(1,845)
Property and equipment, net	<u>\$ 8,486</u>	<u>\$ 5,311</u>

Depreciation is determined using the straight-line method over the estimated useful lives of the respective assets, generally three to five years. Leasehold improvements are amortized on a straight-line basis over the shorter of their estimated useful lives or the term of the lease. Depreciation and amortization expense was \$1.2 million, \$0.8 million and \$0.5 million during the years ended December 31, 2021, 2020 and 2019, respectively.

**Impairment of Long-lived Assets**

Long-lived assets consist primarily of property and equipment and are reviewed for impairment whenever events or changes in circumstances indicate that the carrying amount of an asset may not be recoverable. If circumstances require that an asset be tested for possible impairment, we compare the undiscounted cash flows expected to be generated by the asset to the carrying amount of the asset. If the carrying amount of the asset is not recoverable on an undiscounted cash flow basis, we determine the fair value of the asset and recognize an impairment loss to the extent the carrying amount of the asset exceeds its fair value. We determine fair value using the income approach based on the present value of expected future cash flows or other appropriate measures of estimated fair value. Our cash flow assumptions consider historical and forecasted revenue and operating costs and other relevant factors. We did not record any impairment charges on long-lived assets during the years ended December 31, 2021, 2020, or 2019.

**Accrued Expenses**

Accrued expenses consisted of the following:

	<b>December 31,</b>	
	<b>2021</b>	<b>2020</b>
Payroll related	\$ 17,655	\$ 11,965
Interest	160	160
Product warranty liability	468	159
Current operating lease liability	312	—
Other accrued expenses	1,859	1,232
Total accrued expenses	<u>\$ 20,454</u>	<u>\$ 13,516</u>

**INSPIRE MEDICAL SYSTEMS, INC.**  
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The following table shows the changes in our estimated product warranty liability accrual, included in accrued liabilities:

	Year Ended December 31,		
	2021	2020	2019
Balance at beginning of period	\$ 159	\$ 115	\$ 21
Accruals of warranties issued	576	137	156
Settlements of warranty claims	(267)	(93)	(62)
Balance at the end of the period	<u>\$ 468</u>	<u>\$ 159</u>	<u>\$ 115</u>

### Revenue Recognition

We recognize revenue in accordance with ASC Topic 606, *Revenue from Contracts with Customers* ("ASC 606"). Revenues from product sales are recognized when the customer obtains control of the product, which occurs at a point in time, either upon shipment of the product or receipt of the product, depending on shipment terms. Our standard shipping terms are free on board shipping point, unless the customer requests that control and title to the inventory transfer upon delivery. In those cases where shipping and handling costs are billed to customers, we classify the amounts billed as a component of cost of goods sold.

Revenue is measured as the amount of consideration we expect to receive, adjusted for any applicable estimates of variable consideration and other factors affecting the transaction price, which is based on the invoiced price, in exchange for transferring products. All revenue is recognized when we satisfy our performance obligations under the contract. The majority of our contracts have a single performance obligation and are short term in nature.

Sales taxes and value added taxes in foreign jurisdictions that are collected from customers and remitted to governmental authorities are accounted for on a net basis and therefore are excluded from net sales. Shipping and handling costs associated with outbound freight after control over a product has transferred to a customer are accounted for as a fulfillment cost and are included in cost of goods sold.

Variable consideration related to certain customer sales incentives is estimated based on the amounts expected to be paid based on the agreement with the customer using probability assessments.

We offer customers a limited right of return for our product in case of non-conformity or performance issues. We estimate the amount of our product sales that may be returned by our customers based on historical sales and returns. As our historical product returns to date have been immaterial, we have not recorded a reduction in revenue related to variable consideration for product returns.

See Note 9 for disaggregated revenue by geographic area.

### Cost of Goods Sold

Cost of goods sold consists primarily of acquisition costs for the components of the Inspire system, overhead costs, scrap and inventory obsolescence, warranty replacement costs, as well as distribution-related expenses such as logistics and shipping costs, net of shipping costs charged to customers. The overhead costs include the cost of material procurement, depreciation expense for production equipment, and operations supervision and management personnel, including employee compensation, stock-based compensation, supplies, and travel.

### Research and Development

Research and development expenses consist primarily of product development, clinical and regulatory affairs, quality assurance, consulting services, and other costs associated with products and technologies in development. These expenses include employee compensation, including stock-based compensation, supplies, materials,

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consulting, and travel expenses related to research and development programs. Clinical expenses include clinical trial design, clinical site reimbursement, data management, travel expenses, and the cost of manufacturing products for clinical trials.

### **Stock-Based Compensation**

We maintain an equity incentive plan to provide long-term incentives for eligible employees, consultants, and members of the board of directors. The plan allows for the issuance of restricted stock units ("RSUs") and non-statutory and incentive stock options to employees and RSUs and non-statutory stock options to consultants and directors. We also offer an employee stock purchase plan which allows participating employees to purchase shares of our common stock at a discount through payroll deductions.

We recognize equity-based compensation expense for awards of equity instruments to employees and directors based on the grant date fair value of those awards in accordance with ASC Topic 718, *Stock Compensation* ("ASC 718"). ASC 718 requires all equity-based compensation awards to employees and directors, including grants of RSUs and stock options, to be recognized as expense in the statements of operations and comprehensive loss based on their grant date fair values. We estimate the fair value of stock options using the Black-Scholes option pricing model and the fair value of RSUs is equal to the closing price of our common stock on the grant date. The fair value of each purchase under the employee stock purchase plan is estimated at the beginning of the offering period using the Black-Scholes option pricing model. We have not granted any stock-based awards to our consultants.

The Black-Scholes option pricing model requires the input of certain subjective assumptions, including (i) the expected share price volatility, (ii) the expected term of the award, (iii) the risk-free interest rate and (iv) the expected dividend yield. Due to our limited operating history and a lack of company specific historical and implied volatility data, we have incorporated our historical stock trading volatility with those of a peer group of public companies for the calculation of volatility. The group of peer companies have characteristics similar to us, including stage of product development and focus on the life science industry. The historical volatility is calculated based on a period of time commensurate with the expected term assumption. We use the simplified method, which is the average of the final vesting tranche date and the contractual term, to calculate the expected term for options granted to employees and directors as we do not have sufficient historical exercise data to provide a reasonable basis upon which to estimate the expected term. The risk-free interest rate is based on a U.S. government Treasury instrument whose term is consistent with the expected term of the stock options. We use an assumed dividend yield of zero as we have never paid dividends and have no current plans to pay any dividends on our common stock.

We expense the fair value of our equity-based compensation awards granted to employees and directors on a straight-line basis over the associated service period, which is generally the period in which the related services are received. We account for award forfeitures as they occur.

### **Advertising Expenses**

We expense the costs of advertising, including promotional expenses, as incurred. Advertising expenses were \$47.8 million, \$26.4 million, and \$18.0 million during the years ended December 31, 2021, 2020, and 2019, respectively.

### **Leases**

Operating leases are included in operating lease right-of-use ("ROU") asset, accrued expenses, and operating lease liability – non-current portion in our balance sheets. ROU assets represent our right to use an underlying asset for the lease term and lease liabilities represent our obligation to make lease payments arising from the lease. Operating lease ROU assets and liabilities are recognized at the lease commencement date based on the present value of lease payments over the lease term. In determining the present value of lease payments, we use our incremental borrowing rate based on the information available at the lease commencement date as the rate implicit in the lease is not readily determinable. The determination of our incremental borrowing rate requires management

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judgment based on information available at lease commencement. The operating lease ROU assets also include adjustments for prepayments, accrued lease payments and exclude lease incentives. Our lease terms may include options to extend or terminate the lease when it is reasonably certain that we will exercise such options. Operating lease cost is recognized on a straight-line basis over the expected lease term. Lease agreements entered into after the adoption of ASC 842 that include lease and non-lease components are accounted for as a single lease component. Lease agreements with a noncancelable term of less than 12 months are not recorded on our balance sheets.

#### **Income Taxes**

We account for income taxes using the liability method. Under this method, deferred tax assets and liabilities are determined based on the differences between the financial reporting and tax bases of assets and liabilities and are measured using the enacted tax rates that will be in effect when the differences are expected to reverse. Valuation allowances against deferred tax assets are established, when necessary, to reduce deferred tax assets to the amounts expected to be realized. As we have historically incurred operating losses, we have recorded a full valuation allowance against our net deferred tax assets, and there is no provision for income taxes other than minimal state taxes and an accrual for uncertain tax benefits. Our policy is to record interest and penalties expense related to uncertain tax positions as other expense in the statements of operations and comprehensive loss.

#### **Comprehensive Loss**

Comprehensive loss consists of net loss and changes in unrealized gains and losses due to interest rate fluctuations and other external factors on investments classified as available-for-sale. Accumulated other comprehensive (loss) income is presented in the accompanying balance sheets as a component of stockholders' equity.

#### **Loss Per Share**

Basic net loss per share is computed by dividing the net loss by the weighted average number of shares of common stock outstanding during the period. Diluted net loss per share is computed by dividing the net loss by the weighted average number of shares of common stock and dilutive potential shares of common stock outstanding during the period. Because we have reported a net loss for all periods presented, diluted net loss per share is the same as basic net loss per share as all potentially dilutive shares consisting of outstanding stock options, unvested RSUs, and shares issuable under our employee stock purchase plan were antidilutive in those periods.

#### **Purchase Commitments**

As of December 31, 2021, we had purchase commitments to suppliers for purchases totaling \$114.6 million.

#### **Recent Accounting Pronouncements**

We have reviewed and considered all recent accounting pronouncements that have not yet been adopted and believe there are none that could potentially have a material impact on our business practices, financial condition, results of operations, or disclosures.

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### 3. Investments

Our investments are classified as available-for-sale and consist of the following:

	December 31, 2021			
	Amortized Cost	Unrealized Gross		Aggregate Fair Value
		Gains	Losses	
Long-Term:				
U.S. government securities	\$ 9,993	\$ —	\$ (55)	\$ 9,938
Long-term investments	\$ 9,993	\$ —	\$ (55)	\$ 9,938

	December 31, 2020			
	Amortized Cost	Unrealized Gross		Aggregate Fair Value
		Gains	Losses	
Short-Term:				
Commercial paper	\$ 13,275	\$ —	\$ —	\$ 13,275
Corporate bonds	6,543	—	(3)	6,540
U.S. government securities	23,997	32	—	24,029
Short-term investments	\$ 43,815	\$ 32	\$ (3)	\$ 43,844

As of December 31, 2021 and 2020, we had no investments with a contractual maturity of greater than two years. Currently, we do not intend to sell the investments and it is not more likely than not that we will be required to sell the investments before recovery of their amortized cost bases, which may be maturity. We do not consider those investments to be other-than-temporarily impaired as of December 31, 2021. At the end of each reporting period, we evaluate potential credit impairment on available-for-sale securities in an unrealized loss position, based on the expected cash flows to be collected and the yield-to-maturity on those securities. Securities with a valuation allowance for expected credit losses and deemed uncollectible are permanently written-down, and a reversal out of the valuation allowance occurs.

### 4. Leases

#### Adoption of ASC Topic 842, *Leases*

On December 31, 2019, we adopted the new accounting standard ASC 842, *Leases*, which requires lessees to recognize a lease liability and a ROU asset for all leases with lease terms greater than 12 months. We used the effective date of this standard as the date of initial application, with no retrospective adjustments to prior comparative periods. We were an emerging growth company as defined by the JOBS Act until December 31, 2019 and therefore this guidance became effective for us on the Annual Report on Form 10-K for the fiscal year ended December 31, 2019.

#### Leases Overview

We previously rented approximately 9,500 square feet of office space under an operating lease that expired on March 31, 2019. In September 2018, we entered into a non-cancelable operating lease agreement to sublease approximately 45,000 square feet of office space for our corporate headquarters, which included real estate taxes and operating expenses in the base rent. This lease commenced January 15, 2019 and expired November 30, 2020.

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In May 2019, we entered into a new, non-cancelable operating lease agreement for the same space directly with the landlord. The initial lease term commenced on December 1, 2020 and expires May 31, 2028 with an option to renew for one additional period of five years at the then-prevailing market rate. The exercise of the lease renewal option is at our sole discretion and was not included in the lease term for the calculation of the ROU asset and lease liability when the lease commenced on December 1, 2020 as it is not reasonably certain of exercise.

Beginning December 1, 2020, in addition to base rent, we also pay our proportionate share of the operating expenses, as defined in the lease. These payments are made monthly, and are adjusted annually to reflect actual charges incurred for operating expenses, such as common area maintenance, taxes and insurance. In conjunction with this lease, the landlord provided Inspire with a \$0.6 million rent abatement and a refurbishment allowance in the amount of the cost of any leasehold improvements, not to exceed \$1.1 million upon Inspire providing the necessary documentation evidencing the costs of the leasehold improvements that are completed by May 31, 2022. However, the lease allows us to allocate the refurbishment allowance against base rent instead of taking a tenant improvement reimbursement. At this time, we intend to allocate the full amount of the refurbishment allowance against base rent.

In August 2021, we entered into a new, non-cancelable operating lease agreement for approximately 25,000 square feet of additional corporate office space with our existing landlord. The initial lease term commenced on October 1, 2021 and expires May 31, 2028 with an option to renew for one additional period of five years at the then-prevailing market rate. The exercise of the lease renewal option is at our sole discretion and was not included in the lease term for the calculation of the ROU asset and lease liability when the lease commenced on October 1, 2021 as it is not reasonably certain of exercise.

Beginning October 1, 2021, in addition to base rent on the additional space, we also pay our proportionate share of the operating expenses, as defined in the lease. These payments are made monthly and will be adjusted annually to reflect actual charges incurred for operating expenses, such as common area maintenance, taxes and insurance. In conjunction with this lease, the landlord provided a \$0.2 million rent abatement and a refurbishment allowance in the amount of the cost of any leasehold improvements, not to exceed \$0.4 million upon Inspire providing the necessary documentation evidencing the costs of the leasehold improvements that are completed by May 31, 2022. However, the lease allows us to allocate the refurbishment allowance against base rent instead of taking a tenant improvement reimbursement. At this time, we intend to allocate the full amount of the refurbishment allowance against base rent.

The following table presents the lease balances within the balance sheets:

	<b>December 31,</b>	
	<b>2021</b>	<b>2020</b>
<b>Right-of-use assets:</b>		
Operating lease right-of-use asset	\$ 7,919	\$ 5,805
<b>Operating lease liabilities:</b>		
Accrued liabilities	312	—
Operating lease liability, non-current portion	8,796	5,886
Total operating lease liabilities	<u>\$ 9,108</u>	<u>\$ 5,886</u>

The cost components of our operating leases were as follows:

	<b>Year Ended December 31,</b>		
	<b>2021</b>	<b>2020</b>	<b>2019</b>
Operating lease cost	\$ 1,125	\$ 1,035	\$ 1,021
Variable lease cost	1,001	72	29
Total lease cost	<u>\$ 2,126</u>	<u>\$ 1,107</u>	<u>\$ 1,050</u>



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Variable lease costs consist primarily of taxes, insurance and common area maintenance costs on our current corporate office leases, the first of which commenced December 1, 2020, and our prior corporate office lease, which expired March 31, 2019.

Maturities of our lease liability for our operating lease are as follows as of December 31, 2021:

2022	\$	725
2023		1,777
2024		1,830
2025		1,885
2026		1,941
Thereafter		2,841
Total undiscounted lease payments		10,999
Less: imputed interest		(1,891)
Present value of lease liability	\$	9,108

As of December 31, 2021, the remaining lease term on both operating leases was 6.4 years and the weighted average discount rate was 5.3%. The operating cash outflows from our operating leases were \$0.1 million, \$1.0 million, and \$1.0 million for the years ended December 31, 2021 and 2020, and 2019, respectively.

## 5. Long-Term Debt

### Credit Facility

In August 2015, we entered into a loan and security agreement that initially provided for a term A loan facility in the amount of \$15.5 million, which was fully funded on the closing date, and a term B loan facility in an amount between \$3.5 million and \$10.0 million, subject to our achievement of certain revenue milestones. We refer to our term A loan facility and our term loan B facility together as our credit facility.

In February 2017, we amended the loan and security agreement to, among other things, increase borrowings under the term A loan facility by \$1.0 million, for a total of \$16.5 million outstanding under the credit facility and reduced borrowings available under the term B loan facility to \$9.0 million.

In February 2018, we borrowed an additional \$8.0 million under the term B loan facility portion of the credit facility for a total of \$24.5 million outstanding under the credit facility.

In March 2019, we amended the loan and security agreement. The amendment modified the terms of the loan and security agreement to: (1) extend the interest-only date from March 1, 2020 to April 1, 2022 and extend the maturity date from February 1, 2022 to March 1, 2024; (2) reduce the final payment percentage from 5.50% to 3.50%; (3) modify the basic rate to be a per annum rate of interest (based on a year of 360 days) equal to the sum of (i) the greater of (A) the 30 day U.S. LIBOR rate reported in *The Wall Street Journal* on the last business day of the month that immediately precedes the month in which the interest will accrue or (B) 2.50%, plus (ii) 5.10%; provided, however, under no circumstances will the basic rate be less than 7.60%; (4) provide a mechanism for determining an alternative interest rate to replace the U.S. LIBOR rate upon the occurrence of certain circumstances; and (5) revise the prepayment fee to be between 1.00% and 3.00% of the principal amount, depending on the timing of any prepayment. Upon closing the amendment to the loan and security agreement, payment of the previously accrued final payment under the credit facility was required.

In addition to the principal and interest payments, under the credit facility, we are required to pay a final payment fee of 3.50% on all amounts outstanding, which is being accreted using the effective interest rate method over the term

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of the loan and security agreement and shall be due at the earlier of maturity or prepayment. Borrowings are prepayable at our option in whole, but not in part, together with all accrued and unpaid interest thereon and, if not previously made, the final payment, subject to a prepayment fee of 1.00%.

The credit facility includes affirmative and restrictive covenants and events of default, including the following events of default: payment defaults, breaches of covenants, judgment defaults, cross defaults to certain other contracts, certain events with respect to governmental approvals if such events could cause a material adverse change, a material impairment in the perfection or priority of the lender's security interest or in the value of the collateral, a material adverse change in the business, operations, or condition of us or any of our subsidiaries, and a material impairment of the prospect of repayment of the loans. Upon the occurrence of an event of default, a default increase in the interest rate of an additional 5.00% could be applied to the outstanding loan balance and the lender could declare all outstanding obligations immediately due and payable and take such other actions as set forth in the loan and security agreement.

Our obligations under the credit facility are secured by a first priority security interest in substantially all of our assets, other than our intellectual property. There are no financial covenants contained in the loan and security agreement. We were in compliance with the affirmative and restrictive covenants as of December 31, 2021.

Expected future principal payments for the credit facility are as follows:

**Year ending December 31:**

2022	\$ 9,188
2023	12,250
2024	3,062
Total expected future principal payments	<u>\$ 24,500</u>

## **6. Employee Retirement Plan**

We sponsor a defined contribution employee retirement plan covering all of our full-time employees. The plan allows for eligible employees to defer a portion of their eligible compensation up to the maximum allowed by IRS Regulations. As of December 31, 2021, we had not made contributions since inception. Beginning January 1, 2022, we elected to begin making voluntary matching contributions to the plan. We match 50% of the first 6% of each participating employee's contribution, up to 3% of eligible earnings. Our match contributions are made to funds designated by the participant, none of which are based on Inspire common stock.

## **7. Stockholders' Equity and Stock-Based Compensation**

### **Preferred Stock**

We have 10,000,000 shares of authorized preferred stock. As of December 31, 2020 and 2021, no preferred stock had been issued.

### **Stock-Based Compensation Plans**

We adopted the 2007 Stock Incentive Plan (the "2007 Plan") in November 2007, which terminated in accordance with its terms on November 28, 2017; however, the outstanding stock options may continue to be exercised in accordance with their terms. Immediately following the termination of the 2007 Plan, we adopted the 2017 Stock Incentive Plan (the "2017 Plan"), which contains substantially similar terms and conditions as the 2007 Plan. Upon the IPO, no further grants were made under the 2017 Plan and we adopted the 2018 Stock Incentive Plan (the "2018 Plan").

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The purpose of the 2018 Plan is to promote the interest of our company and our stockholders by aiding in attracting and retaining employees, officers, consultants, independent contractors, and directors capable of assuring the future success of our business and to afford such persons an opportunity to acquire a proprietary interest in our company. The board of directors may amend, alter, suspend, discontinue, or terminate the 2018 Plan at any time with the approval of our stockholders. A total of 1,386,809 shares of common stock were initially reserved for issuance under the 2018 Plan, and this share reserve will automatically be supplemented each January 1, commencing on January 1, 2019 and ending on and including January 1, 2028, by an amount of shares equal to the lesser of: a) 739,631 shares, b) 4% of the shares outstanding on the final day of the immediately preceding fiscal year and c) such smaller number of shares as determined by the board of directors. As of December 31, 2021, there were 3,250,937 shares reserved for issuance under the 2018 Plan, of which 1,272,049 shares were available for issuance.

The following table presents the components and classification of stock-based compensation expense recognized for stock options and RSUs granted under the 2017 Plan, the 2018 Plan and under our Employee Stock Purchase Plan (the "ESPP"):

	Year Ended December 31,		
	2021	2020	2019
Stock options	\$ 24,703	\$ 11,823	\$ 5,699
Restricted stock units	105	—	—
Employee stock purchase plan	1,370	994	498
Total stock-based compensation	<u>\$ 26,178</u>	<u>\$ 12,817</u>	<u>\$ 6,197</u>
Cost of goods sold	\$ 341	\$ 170	\$ 121
Research and development	4,419	1,931	689
Selling, general and administrative	21,418	10,716	5,387
Total stock-based compensation	<u>\$ 26,178</u>	<u>\$ 12,817</u>	<u>\$ 6,197</u>

### Stock Options

Options are granted at the exercise price, which is equal to the closing price of our stock on the date of grant. The stock options granted to employees include a four-year service period and 25% vest after the first year of service and the remainder vest in equal monthly installments over the next 36 months of service. The stock options granted to the board of directors vest in one or three equal annual installments, in each case subject to the director's continuous services through the applicable vesting date. The stock options have a contractual life of ten years.

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A summary of stock option activity and related information is as follows:

	<b>Options</b>	<b>Weighted Average Exercise Price</b>	<b>Weighted average remaining contractual term (years)</b>	<b>Aggregate intrinsic value (in thousands)</b>
Outstanding at December 31, 2018	2,745,156	\$ 12.64	7.4	\$ 81,453
Granted	809,454	\$ 67.43		
Exercised	(664,617)	\$ 1.71		\$ 35,305
Forfeited/expired	(45,829)	\$ 36.39		
Outstanding at December 31, 2019	2,844,164	\$ 30.41	7.9	\$ 124,585
Granted	703,838	\$159.67		
Exercised	(624,315)	\$ 11.29		\$ 54,821
Forfeited/expired	(66,123)	\$ 44.57		
Outstanding at December 31, 2020	2,857,564	\$ 66.09	7.9	\$ 351,626
Granted	228,302	\$215.34		
Exercised	(323,860)	\$ 35.44		\$ 58,360
Forfeited/expired	(115,771)	\$118.85		
Outstanding at December 31, 2021	2,646,235	\$ 80.41	7.1	\$ 397,015
Exercisable at December 31, 2021	1,518,096	\$ 41.97	6.2	\$ 285,546

The aggregate intrinsic value of options exercised is the difference between the estimated fair market value of our common stock at the date of exercise and the exercise price for those options. The aggregate intrinsic value of outstanding options is the difference between the closing price as of the date outstanding and the exercise price of the underlying stock options. The total grant date fair value of options vested during the year was \$23.9 million, \$11.1 million and \$5.1 million for the years ended December 31, 2021, 2020 and 2019, respectively.

As of December 31, 2021, the amount of unearned stock-based compensation currently estimated to be expensed from now through the year 2025 related to unvested employee and non-employee director stock options is \$67.5 million which we expect to recognize over a weighted average period of 2.4 years. If there are any modifications or cancellations of the underlying unvested securities, we may be required to accelerate, increase, or cancel any remaining unearned stock compensation expense. Future stock-based compensation expense and unearned stock-based compensation will increase to the extent that we grant additional stock-based awards.

We estimate the fair value of stock-based awards on the date of grant using the Black-Scholes option pricing model using the fair market value of our common stock on the date of grant and a number of other assumptions. These assumptions include estimates regarding the expected term of the awards, estimates of the stock volatility over a duration that approximates the expected term of the awards, estimates of the risk-free rate, and estimates of expected dividend rates.

The fair value of options granted to employees and directors during the years ended December 31, 2021, 2020, and 2019 was estimated as of the grant date using the Black-Scholes option pricing model using the following assumptions:

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	Year Ended December 31,		
	2021	2020	2019
Expected term (years)	5.50 - 6.25	5.50 - 6.25	5.50 - 6.25
Expected volatility	54.9 - 55.9%	42.3 - 49.3%	43.1 - 50.6%
Risk-free interest rate	0.79 - 1.44%	0.36 - 1.42%	1.39 - 2.63%
Dividend yield	—%	—%	—%
Weighted average fair value	\$113.71	\$73.93	\$30.69

*Expected Term* — Due to our limited amount of historical exercise, forfeiture, and expiration activity, we have opted to use the "simplified method" for estimating the expected term of options, whereby the expected term equals the arithmetic average of the vesting terms and the original contractual term of the option. We will continue to analyze our expected term assumption as more historical data becomes available.

*Expected Volatility* — Due to our limited operating history and a lack of company specific historical and implied volatility data, we have incorporated our historical stock trading volatility with those of a group of similar companies that are publicly traded for the calculation of volatility. When selecting this peer group of public companies on which we have based our expected stock price volatility, we generally selected companies with comparable characteristics to it, including enterprise value, stages of clinical development, risk profiles, position within the industry, and with historical share price information sufficient to meet the expected life of the stock-based awards. We will continue to analyze the historical stock price volatility assumption as more historical data for our common stock becomes available.

*Risk-Free Interest Rate* — The risk-free rate assumption is based on the U.S. government Treasury instruments with maturities similar to the expected term of our stock options.

*Expected Dividend Yield* — The expected dividend assumption is based on our history of not paying dividends and our expectation that we will not declare dividends for the foreseeable future.

The amount of stock-based compensation expense is recognized on a straight-line basis over the vesting term and is reduced by actual forfeitures as they occur.

### Restricted Stock Units

RSUs are share awards that entitle the holder to receive freely tradable shares of our common stock upon vesting. The RSUs cannot be transferred and the awards are subject to forfeiture if the holder's employment terminates prior to the release of the vesting restrictions. The RSUs include a four-year service period and vest in equal installments on each of the first four anniversaries of the date of grant, provided the employee remains continuously employed with the Company. The fair value of the RSUs is equal to the closing price of our common stock on the grant date.

A summary of RSUs and related information is as follows:

	Restricted Stock Units	Weighted Average Grant Date Fair Value	Aggregate Intrinsic Value (in thousands)
Unvested at December 31, 2020	—	\$ —	\$ —
Granted	2,275	\$ 201.51	
Unvested at December 31, 2021	2,275	\$ 201.51	\$ 524

There were no RSUs granted prior to 2021. The aggregate intrinsic value of RSUs outstanding was based on our closing stock price on the last trading day of the period. As of December 31, 2021, there was \$0.4 million of unrecognized stock-based compensation expense related to RSUs to be recognized over a period of 3.1 years.

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**Employee Stock Purchase Plan**

Our employee stock purchase plan ("ESPP") allows participating employees to purchase shares of our common stock at a discount through payroll deductions. The plan is available to all of our U.S.-based full-time employees. Participating employees may purchase common stock, on a voluntary after-tax basis, at a price equal to 85% of the lower of the closing market price per share of our common stock on the first or last trading day of each stock purchase period. The plan provides for six-month purchase periods, beginning on January 1 and July 1 of each calendar year.

A total of 277,362 shares of common stock were initially reserved for issuance under the ESPP, and this share reserve will automatically be supplemented each January 1, commencing on January 1, 2019 and ending on and including January 1, 2028, by an amount of shares equal to the lesser of: a) 184,908 shares, b) 1% of the shares outstanding on the final day of the immediately preceding calendar year and c) such smaller number of shares as the board of directors may determine. During the year ended December 31, 2021, 21,507 shares were purchased under the ESPP, utilizing \$3.5 million of employee contributions. As of December 31, 2021, 744,596 shares were available for future issuance under the ESPP.

**8. Income Taxes**

Due to our cumulative net loss position, a valuation allowance has been required to be established for all deferred tax assets as of December 31, 2021, 2020, and 2019.

The components of our provision for income taxes are as follows:

	December 31,		
	2021	2020	2019
Current			
United States	\$ 23	\$ 70	\$ —
Foreign	49	45	40
Total current	72	115	40
Total provision for income taxes	<u>\$ 72</u>	<u>\$ 115</u>	<u>\$ 40</u>

The reconciliation of taxes at the federal statutory rate to our provision for income taxes are as follows:

	Year Ended December 31,		
	2021	2020	2019
Tax at federal statutory rate	21.0 %	21.0 %	21.0 %
State, net of federal benefit	4.0	3.6	4.3
Stock-based compensation	18.1	8.0	3.9
Research and development ("R&D") tax credit	3.3	1.4	1.0
Other	0.4	(0.1)	(0.6)
Change in valuation allowance	(47.0)	(34.1)	(29.7)
Total	<u>(0.2)%</u>	<u>(0.2)%</u>	<u>(0.1)%</u>

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
(Table amounts in thousands, except share and per share amounts)

Significant components of net deferred tax assets and liabilities were as follows:

	<b>Year Ended December 31,</b>	
	<b>2021</b>	<b>2020</b>
Deferred tax assets:		
Net operating losses	\$ 71,445	\$ 58,150
R&D tax credits	6,257	3,761
R&D expenditures, capitalized for tax	1,505	1,846
Interest expense carryforward	757	252
Accruals and other	2,092	1,200
Depreciation and amortization	62	167
Lease asset	2,277	1,444
Stock-based compensation	4,543	1,770
Other comprehensive loss	14	—
Total deferred tax assets	88,952	68,590
Deferred tax liabilities:		
Depreciation	(59)	—
Lease liability	(1,980)	(1,424)
Other comprehensive income	—	(7)
Total deferred tax liabilities	(2,039)	(1,431)
Net deferred tax assets	86,913	67,159
Valuation allowance	(86,913)	(67,159)
	\$ —	\$ —

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

As of December 31, 2021, our gross federal net operating loss carryforwards of \$286.3 million will expire at various dates beginning in 2028. In addition, net operating loss carryforwards for state income tax purposes of \$198.2 million will begin to expire in 2023. We also have gross R&D credit carryforwards of \$6.7 million as of December 31, 2021 which will expire at various dates beginning in 2033.

Utilization of the net operating loss carryforwards and R&D credit carryforwards may be subject to an annual limitation due to the ownership change limitations provided by Section 382 and Section 383 of the Internal Revenue Code of 1986 and similar state provisions. We have not performed a detailed analysis to determine whether an ownership change has occurred. Such a change of ownership would limit our utilization of the net operating losses and R&D credits and could be triggered by subsequent sales of securities by us or our stockholders.

Realization of the deferred tax assets is dependent upon the generation of future taxable income, if any, the amount and timing of which are uncertain. Based on available objective evidence and cumulative losses, we believe it is more likely than not that the deferred tax assets are not recognizable and will not be recognizable until we have sufficient taxable income. Accordingly, the net deferred tax assets have been fully offset by a valuation allowance. The valuation allowance increased by \$19.8 million and \$19.5 million during the years ended December 31, 2021 and 2020, respectively.

**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
(Table amounts in thousands, except share and per share amounts)

The changes to our gross unrecognized tax benefits were as follows:

	Year Ended December 31,		
	2021	2020	2019
Balance beginning of the year	\$ 85	\$ 40	\$ —
Increase in balances related to current year tax positions	49	—	40
Increase in balances related to prior year tax positions	—	45	—
Balance end of the year	<u>\$ 134</u>	<u>\$ 85</u>	<u>\$ 40</u>

We file income tax returns in the applicable jurisdictions. The 2018 to 2020 tax years remain open to examination by the major taxing authorities to which we are subject. We do not expect a significant change to our unrecognized tax benefits over the next 12 months.

Our policy is to record interest related to uncertain tax positions as interest expense and any penalties as other expense in our statements of operations and comprehensive loss. There was no interest or penalties accrued at December 31, 2021 and 2020.

## 9. Segment Reporting and Revenue Disaggregation

We operate our business as one operating segment. An operating segment is defined as a component of an enterprise for which separate discrete financial information is available and evaluated regularly by the chief operating decision maker, or decision-making group, in deciding how to allocate resources and in assessing performance. Segment information is consistent with how management reviews the business, makes investing and resource allocation decisions and assesses operating performance.

We sell our Inspire system to hospitals and ambulatory surgery centers in the U.S. and in select countries in Europe through a direct sales organization, and in Japan through a distributor. Revenue by geographic region is as follows:

	Year Ended December 31,		
	2021	2020	2019
United States	\$ 220,976	\$ 106,108	\$ 73,660
All other countries	12,418	9,273	8,390
Total revenue	<u>\$ 233,394</u>	<u>\$ 115,381</u>	<u>\$ 82,050</u>

All of our long-lived assets are located in the U.S.

## 10. Loss Per Share

Basic net loss per share is computed by dividing the net loss by the weighted average number of shares of common stock outstanding during the period. Diluted net loss per share is computed by dividing the net loss by the weighted average number of shares of common stock and dilutive potential shares of common stock outstanding during the period. Because we have reported a net loss for all periods presented, diluted net loss per share is the same as basic net loss per share as all of the following potentially dilutive shares were antidilutive in those periods.



**INSPIRE MEDICAL SYSTEMS, INC.**  
**NOTES TO FINANCIAL STATEMENTS**  
**(Table amounts in thousands, except share and per share amounts)**

The following common stock-based awards were excluded from the computation of diluted net loss per common share for the periods presented because including them would have been anti-dilutive:

	<b>Year Ended December 31,</b>		
	<b>2021</b>	<b>2020</b>	<b>2019</b>
Common stock options outstanding	2,646,235	2,857,564	2,844,164
Unvested restricted stock units	2,275	—	—
<b>Total</b>	<b>2,648,510</b>	<b>2,857,564</b>	<b>2,844,164</b>

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

None.

**Item 9A. Controls and Procedures.*****Evaluation of Disclosure Controls and Procedures***

Our management, with the participation of our Chief Executive Officer and Chief Financial Officer, evaluated the effectiveness of our disclosure controls and procedures as of December 31, 2021. The term “disclosure controls and procedures,” as defined in Rules 13a-15(e) and 15d-15(e) under the Exchange Act, means controls and other procedures of a company that are designed to ensure that information required to be disclosed by a company in the reports that it files or submits under the Exchange Act is recorded, processed, summarized and reported, within the time periods specified in the SEC’s rules and forms. Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information required to be disclosed by a company in the reports that it files or submits under the Exchange Act is accumulated and communicated to the company’s management, including its principal executive and principal financial officers, or persons performing similar functions, as appropriate to allow timely decisions regarding required disclosure. Management recognizes that any controls and procedures, no matter how well designed and operated, can provide only reasonable assurance of achieving their objectives and management necessarily applies its judgment in evaluating the cost-benefit relationship of possible controls and procedures. Based on the evaluation of our disclosure controls and procedures as of December 31, 2021, our Chief Executive Officer and Chief Financial Officer concluded that, as of such date, our disclosure controls and procedures were effective at the reasonable assurance level.

***Changes in Internal Control over Financial Reporting***

There was no change in the Company’s internal control over financial reporting, as defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act, that occurred during the quarter ended December 31, 2021 that has materially affected, or is reasonably likely to materially affect, our internal control over financial reporting.

***Management Report on Internal Control over Financial Reporting***

Management is responsible for establishing and maintaining adequate internal control over financial reporting, as such term is defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act. Internal control over financial reporting is a process designed by, or under the supervision of, a company’s principal executive and principal financial officers, or persons performing similar functions, and effected by the company’s board of directors, management and other personnel, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with U.S. generally accepted accounting principles. It includes those policies and procedures that:

- pertain to the maintenance of records that in reasonable detail accurately and fairly reflect the transactions and dispositions of the assets of the company;
- provide reasonable assurance that transactions are recorded as necessary to permit preparation of financial statements in accordance with U.S. generally accepted accounting principles, and that receipts and expenditures of the company are being made only in accordance with authorizations of management and directors of the company; and
- provide reasonable assurance regarding prevention or timely detection of unauthorized acquisition, use or disposition of the company’s assets that could have a material effect on the financial statements.

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

Our management conducted an assessment of the effectiveness of our internal control over financial reporting based on the criteria set forth in “Internal Control-Integrated Framework (2013)” issued by the Committee of Sponsoring Organizations of the Treadway Commission. Based on this assessment, management concluded that, as of December 31, 2021, our internal control over financial reporting was effective.

Ernst & Young LLP, our independent registered public accounting firm that audited the consolidated financial statements included in this Annual Report on Form 10-K, has issued an audit report on our internal control over financial reporting. See Report of Independent Registered Public Accounting Firm below.

## **Report of Independent Registered Public Accounting Firm**

To the Stockholders and the Board of Directors of Inspire Medical Systems, Inc.

### **Opinion on Internal Control Over Financial Reporting**

We have audited Inspire Medical Systems, Inc.'s internal control over financial reporting as of December 31, 2021, based on criteria established in Internal Control—Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework) (the COSO criteria). In our opinion, Inspire Medical Systems, Inc. (the Company) maintained, in all material respects, effective internal control over financial reporting as of December 31, 2021, based on the COSO criteria.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States) (PCAOB), the balance sheets of Inspire Medical Systems, Inc. as of December 31, 2021 and 2020, the related statements of operations and comprehensive loss, stockholders' equity and cash flows for each of the three years in the period ended December 31, 2021, and the related notes and our report dated February 15, 2022 expressed an unqualified opinion thereon.

### **Basis for Opinion**

The Company's management is responsible for maintaining effective internal control over financial reporting and for its assessment of the effectiveness of internal control over financial reporting included in the accompanying Management Report on Internal Control over Financial Reporting. Our responsibility is to express an opinion on the Company's internal control over financial reporting based on our audit. We are a public accounting firm registered with the PCAOB and are required to be independent with respect to the Company in accordance with the U.S. federal securities laws and the applicable rules and regulations of the Securities and Exchange Commission and the PCAOB.

We conducted our audit in accordance with the standards of the PCAOB. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether effective internal control over financial reporting was maintained in all material respects.

Our audit included obtaining an understanding of internal control over financial reporting, assessing the risk that a material weakness exists, testing and evaluating the design and operating effectiveness of internal control based on the assessed risk, and performing such other procedures as we considered necessary in the circumstances. We believe that our audit provides a reasonable basis for our opinion.

### **Definition and Limitations of Internal Control Over Financial Reporting**

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

Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. Also, projections of any evaluation of effectiveness to future periods are subject to the risk

that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

/s/Ernst & Young LLP

Minneapolis, Minnesota

February 15, 2022

**Item 9B. Other Information.**

None.

**Item 9C. Disclosure Regarding Foreign Jurisdictions that Prevent Inspections.**

None.

### PART III

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

The information required by this Item is incorporated by reference to our definitive proxy statement for our 2022 annual meeting of stockholders to be filed with the SEC within 120 days of the fiscal year ended December 31, 2021.

**Item 11. Executive Compensation.**

The information required by this Item is incorporated by reference to our definitive proxy statement for our 2022 annual meeting of stockholders to be filed with the SEC within 120 days of the fiscal year ended December 31, 2021.

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

The following table provides information as of December 31, 2021, regarding our common stock that may be issued under the Inspire Medical Systems, Inc. 2007 Incentive Award Plan, as amended (the "2007 Plan"), the Inspire Medical Systems, Inc. 2017 Incentive Award Plan, as amended (the "2017 Plan"), the Inspire Medical Systems, Inc. 2018 Incentive Award Plan (the "2018 Plan") and the Inspire Medical Systems, Inc. 2018 Employee Stock Purchase Plan (the "2018 ESPP").

Plan category:	Number of Securities to be Issued Upon Exercise of Outstanding Options and Restricted Stock Units	Weighted-Average Exercise Price of Outstanding Options	Number of Securities Available for Future Issuance Under Equity Compensation Plans (excludes securities reflected in column (a))
(a)	(b)	(c)	
Equity compensation plans approved by stockholders			
2007 Plan (1)	524,026	\$ 1.46	—
2017 Plan (1)	145,596	\$ 9.59	—
2018 Plan (2)	1,978,888	\$ 106.56	1,272,049
2018 ESPP (3)	—	\$ —	744,596
Equity compensation plans not approved by	—	\$ —	—
<b>Total</b>	<b><u>2,648,510</u></b>	<b><u>\$ 80.41</u></b>	<b><u>2,016,645</u></b>

(1) The 2007 Plan terminated in accordance with its terms on November 28, 2017; however, outstanding stock options may continue to be exercised in accordance with their terms. In connection with our IPO, we adopted the 2018 Plan and do not make grants or awards under the 2017 Plan.

(2) Pursuant to the terms of the 2018 Plan, the number of shares of common stock available for issuance under the 2018 Plan automatically increases on each January 1, until and including January 1, 2028, by an amount equal to the lesser of (a) 739,631 shares, (b) 4% of the number of shares of common stock outstanding (on an as-converted basis) on the last day of the immediately preceding fiscal year, and (c) such smaller number of shares as is determined by our board of directors. The weighted average exercise price is calculated without taking into account restricted stock that will become issuable, without any cash consideration or other payment, as vesting requirements are achieved.

(3) Pursuant to the terms of the 2018 ESPP Plan, the number of shares reserved under the 2018 ESPP Plan will automatically be supplemented each January 1, until and including January 1, 2028, by an amount of shares equal to the lesser of a) 184,908 shares, b) 1% of the shares outstanding on the final day of the immediately preceding calendar year, and c) such smaller number of shares as the board of directors may determine.

#### **Other**

The remaining information required by this Item is incorporated by reference to our definitive proxy statement for our 2022 annual meeting of stockholders to be filed with the SEC within 120 days of the fiscal year ended December 31, 2021.

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

The information required by this Item is incorporated by reference to our definitive proxy statement for our 2022 annual meeting of stockholders to be filed with the SEC within 120 days of the fiscal year ended December 31, 2021.

#### **Item 14. Principal Accounting Fees and Services.**

Our independent registered public accounting firm is Ernst & Young LLP, Minneapolis, MN, Auditor Firm ID: 42.

The information required by this Item is incorporated by reference to our definitive proxy statement for our 2022 annual meeting of stockholders to be filed with the SEC within 120 days of the fiscal year ended December 31, 2021.

## PART IV

### Item 15. Exhibits, Financial Statement Schedules.

(a) List of documents filed as part of this Annual Report on Form 10-K:

(1) Financial Statements

The financial statements included in Part II, Item 8 of this document are filed as part of this Annual Report on Form 10-K.

(2) Financial Statement Schedules

All schedules are omitted because they are not applicable or the amounts are immaterial or the required information is presented in the consolidated financial statements and notes thereto in Part II, Item 8 above.

(3) Exhibits

The following documents are filed as exhibits to this Annual Report on Form 10-K.

Exhibit Number	Description	Form	File No.	Exhibit	Filing Date	Filed/ Furnished Herewith
3.1	Seventh Amended and Restated Certificate of Incorporation of Inspire Medical Systems, Inc.	8-K	001-38468	3.1	5/7/2018	
3.2	Amended and Restated Bylaws of Inspire Medical Systems, Inc.	8-K	001-38468	3.2	5/7/2018	
4.1	Form of Certificate of Common Stock	S-1	333-224176	4.1	4/23/2018	
4.2	Fifth Amended and Restated Investor Rights Agreement, dated October 25, 2016, among Inspire Medical Systems, Inc. and the Investors party thereto	S-1	333-224176	4.2	4/6/2018	
4.3	Amendment No. 1 to Fifth Amended and Restated Investor Rights Agreement, dated April 20, 2018, among Inspire Medical Systems, Inc. and the Investors party thereto	S-1	333-224176	4.3	4/23/2018	
4.6	Description of Capital Stock	10-K	001-38468	4.6	2/25/2020	
10.1	Assignment and License Agreement, dated as of November 28, 2007, by and between Inspire Medical Systems, Inc. and Medtronic, Inc.	S-1	333-224176	10.1	4/6/2018	
10.2	First Amendment to Assignment and License Agreement, dated as of February 4, 2010, by and between Inspire Medical Systems, Inc. and Medtronic, Inc.	S-1	333-224176	10.2	4/6/2018	
10.3	Loan and Security Agreement, dated as of August 7, 2015, by and between Inspire Medical Systems, Inc. and Oxford Finance LLC	S-1	333-224176	10.3	4/6/2018	
10.4	First Amendment to Loan and Security Agreement, dated as of February 24, 2017, by and between Inspire Medical Systems, Inc. and Oxford Finance LLC	S-1	333-224176	10.4	4/6/2018	
10.5	Second Amendment to the Loan and Security Agreement, dated as of March 27, 2019, by and between Inspire Medical Systems, Inc. and Oxford Finance LLC.	8-K	001-38468	10.1	3/29/2019	
10.6†	2007 Stock Incentive Plan, as amended	S-1	333-224176	10.6	4/6/2018	
10.7†	Form of Incentive Stock Option Agreement pursuant to 2007 Stock Incentive Plan	S-1	333-224176	10.7	4/6/2018	
10.8†	2017 Stock Incentive Plan, as amended	S-1	333-224176	10.8	4/6/2018	



Exhibit Number	Description	Form	File No.	Exhibit	Filing Date	Filed/ Furnished Herewith
10.9†	Form of Incentive Stock Option Agreement pursuant to 2017 Stock Incentive Plan	S-1	333-224176	10.9	4/6/2018	
10.10†	Form of Non-Statutory Stock Option Agreement pursuant to 2017 Stock Incentive Plan	S-1	333-224176	10.10	4/6/2018	
10.11†	Inspire Medical Systems, Inc. 2018 Incentive Award Plan	S-1	333-224176	10.11	4/23/2018	
10.12†	Form of Stock Option Award Agreement under Inspire Medical Systems, Inc. 2018 Incentive Award Plan	S-1	333-224176	10.12	4/23/2018	
10.13†	Form of Restricted Stock Unit Award Agreement under Inspire Medical Systems, Inc. 2018 Incentive Award Plan	10-K	001-38468	10.13	2/24/2021	
10.14†	Form of Performance Stock Unit Award Agreement under Inspire Medical Systems, Inc. 2018 Incentive Award Plan					*
10.15†	Inspire Medical Systems, Inc. 2018 Employee Stock Purchase Plan	S-1	333-224176	10.13	4/23/2018	
10.16†	Amended and Restated Employment Agreement, by and between Inspire Medical Systems, Inc. and Timothy Herbert	S-1	333-224176	10.15	4/23/2018	
10.17†	Amended and Restated Employment Agreement, by and between Inspire Medical Systems, Inc. and Randy Ban	S-1	333-224176	10.17	4/23/2018	
10.18†	Amended and Restated Employment Agreement, by and between Inspire Medical Systems, Inc. and Richard Buchholz	S-1	333-224176	10.19	4/23/2018	
10.19†	Amended and Restated Employment Agreement, by and between Inspire Medical Systems, Inc. and Steven Jandrich	S-1	333-224176	10.21	4/23/2018	
10.20†	Employment Agreement between the Company and Philip Ebeling	8-K	001-38468	10.1	6/8/2020	
10.21†	Employment Agreement between the Company and Bryan Phillips	10-K	001-38468	10.20	2/24/2021	
10.22†	Form of Indemnification Agreement between Inspire Medical Systems, Inc. and its directors and officers	S-1	333-224176	10.23	4/6/2018	
10.23†	Inspire Medical Systems, Inc. Non-Employee Director Compensation Policy	10-Q	001-38468	10.1	8/3/2021	
21.1	Subsidiaries of Inspire Medical Systems, Inc.					*
23.1	Consent of Ernst & Young LLP					*
31.1	Certification of Principal Executive Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002					*
31.2	Certification of Principal Financial Officer Pursuant to Rules 13a-14(a) and 15d-14(a) under the Securities Exchange Act of 1934, as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002					*
32.1	Certification of Principal Executive Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002					**
32.2	Certification of Principal Financial Officer Pursuant to 18 U.S.C. Section 1350, as Adopted Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002					**
101.INS	Inline XBRL Instance Document - the instance document does not appear in the Interactive Data File because its XBRL tags are embedded within the Inline XBRL document.					*

Exhibit Number	Description	Form	File No.	Exhibit	Filing Date	Filed/ Furnished Herewith
101.SCH	Inline XBRL Taxonomy Extension Schema Document					*
101.CAL	Inline XBRL Taxonomy Extension Calculation Linkbase Document					*
101.DEF	Inline XBRL Taxonomy Extension Definition Linkbase Document					*
101.LAB	Inline XBRL Taxonomy Extension Label Linkbase Document					*
101.PRE	Inline XBRL Taxonomy Extension Presentation Linkbase Document					*
104	Cover Page Interactive Data File (formatted as Inline XBRL and contained in Exhibit 101)					*

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\* Filed herewith.

\*\* Furnished herewith.

† Denotes a management contract or compensation plan or arrangement.

Certain agreements filed as exhibits to this Annual Report on Form 10-K contain representations and warranties that the parties thereto made to each other. These representations and warranties have been made solely for the benefit of the other parties to such agreements and may have been qualified by certain information that has been disclosed to the other parties to such agreements and that may not be reflected in such agreements. In addition, these representations and warranties may be intended as a way of allocating risks among parties if the statements contained therein prove to be incorrect, rather than as actual statements of fact. Accordingly, there can be no reliance on any such representations and warranties as characterizations of the actual state of facts. Moreover, information concerning the subject matter of any such representations and warranties may have changed since the date of such agreements.

#### Item 16. Form 10-K Summary.

None.

## 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.

**Inspire Medical Systems, Inc.**

Date: February 15, 2022

By: /s/ TIMOTHY P. HERBERT

Timothy P. Herbert  
President and Chief Executive Officer  
(principal executive 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 indicated on February 15, 2022.

Signature	Title
<u>/s/ TIMOTHY P. HERBERT</u> Timothy P. Herbert	Chief Executive Officer (principal executive officer), President and Director
<u>/s/ RICHARD BUCHHOLZ</u> Richard Buchholz	Chief Financial Officer (principal financial and accounting officer)
<u>/s/ MARILYN CARLSON NELSON</u> Marilyn Carlson Nelson	Chair of the Board of Directors
<u>/s/ SHELLEY BROADER</u> Shelly Broader	Director
<u>/s/ GARY L. ELLIS</u> Gary L. Ellis	Director
<u>/s/ JERRY GRIFFIN, M.D.</u> Jerry Griffin, M.D.	Director
<u>/s/ DANA G. MEAD, JR.</u> Dana G. Mead, Jr.	Director
<u>/s/ GEORGIA MELENIKIOTOU</u> Georgia Melenikiotou	Director
<u>/s/ SHAWN T MCCORMICK</u> Shawn T McCormick	Director
<u>/s/ CASEY TANSEY</u> Casey Tansey	Director

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## Board of Directors

**Marilyn Carlson Nelson**  
Chair of the Board, Former  
Co-Chair & Co-Chief  
Executive Officer, Carlson  
Holdings, Inc.

**Timothy P. Herbert**  
President and Chief  
Executive Officer, Inspire  
Medical Systems, Inc.

**Shelley Broader**  
Former Chief Executive  
Officer, Chico's FAS

**Gary L. Ellis**  
Former Chief Financial  
Officer, Medtronic, Inc.

**Georgia Garinois-  
Melenikiotou**  
Former Executive Vice  
President, The Estée  
Lauder Companies

**Jerry Griffin, M.D.**  
President, Griffin &  
Schwartz, Scientific  
Services, Inc.

**Shawn T McCormick**  
Former Chief Financial  
Officer, Aldevron, LLC  
& Tornier N.V.

**Dana G. Mead, Jr.**  
Former Chief Executive  
Officer, HeartFlow, Inc.

**Casey Tansey**  
General Partner, U.S.  
Venture Partners

## Senior Leadership

**Timothy P. Herbert**  
President and Chief  
Executive Officer

**Richard J. Buchholz**  
Chief Financial Officer

**Randall A. Ban**  
Chief Commercial Officer

**Philip J. Ebeling**  
Chief Operating Officer

**Bryan K. Phillips**  
Chief Compliance Officer,  
General Counsel,  
and Secretary

**Steven L. Jandrich**  
Vice President,  
Human Resources

**Andreas Henke**  
Senior Vice President,  
European Operations

**Ivan Lubogo**  
Senior Vice President,  
U.S. Sales

**John C. Rondoni**  
Senior Vice President,  
Research, Development  
and Ventures

**Kathy L. Sherwood**  
Senior Vice President,  
Global Market Access

**Martin A. Abrams**  
Vice President, Marketing  
and Customer Experience

## Corporate Information

**Annual Meeting of  
Stockholders**  
Our annual meeting will  
be a completely virtual  
meeting of stockholders.

April 28, 2022 at 8:00 AM CT  
[www.virtualshareholdermeeting.com/INSP2022](http://www.virtualshareholdermeeting.com/INSP2022)

**Stock Exchange**  
New York Stock Exchange  
Symbol: INSP

**Independent Registered  
Public Accounting Firm**  
Ernst & Young LLP  
Minneapolis, Minnesota

**Outside Counsel**  
Stinson Leonard Street LLP  
Minneapolis, Minnesota

Latham & Watkins LLP  
New York, New York

**Investor Inquiries**  
LifeSci Advisors  
[inspire@lifesciadvisors.com](mailto:inspire@lifesciadvisors.com)  
(646) 876-3647

**Transfer Agent & Registrar**  
American Stock Transfer &  
Trust Company, LLC

6201 15th Avenue  
Brooklyn, NY 11219  
(800) 937-5449  
[help@astfinancial.com](mailto:help@astfinancial.com)  
[www.astfinancial.com](http://www.astfinancial.com)

The Inspire Logo and Inspire Cloud are registered trademarks of Inspire Medical Systems, Inc. Bluetooth is a registered trademark of Bluetooth SIG, Inc.

### Safe Harbor Statement

Statements in this document regarding future events and expectations, such as forecasts, plans, trends and projections relating to the Company's business and financial performance, are forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. These forward-looking statements speak only as of the date such statements are made and are subject to risks and uncertainties that could cause the Company's results to differ materially from these statements. These risks and uncertainties are described in the Company's Annual Report on Form 10-K, and Inspire undertakes no obligation to update them unless otherwise required by law.

**No mask.  
No hose.  
Just sleep.**



**Corporate  
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