UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

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

Date of report (Date of earliest event reported): October 8, 2020

SI-BONE, INC.

(Exact name of registrant as specified in its charter)

Delaware

001-38701

26-2216351

(State or other jurisdiction of incorporation or organization)

(Commission File Number) (I.R.S. Employer

Identification No.)

471 El Camino Real Suite 101

Santa Clara, CA 95050 (Address of principal executive offices) (Zip Code)

(408) 207-0700

(Registrant's telephone number, include area code)

N/A

(Former Name or Former Address, if Changed Since Last Report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
 Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
 Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
 Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
 Securities registered pursuant to Section 12(b) of the Act:

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company \square

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.

Item 2.02 – Results of Operations and Financial Condition.

On October 8, 2020, SI-BONE, Inc. (the "Company") issued a press release (the "Press Release") announcing preliminary unaudited revenue for the three and nine months ended September 30, 2020 and providing full year 2020 revenue guidance. A copy of the Press Release is attached as Exhibit 99.1 to this current report on Form 8-K and is incorporated by reference herein.

The information in this Item 2.02 and Exhibit 99.1 of this Current Report on Form 8-K are being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934 as amended (Exchange Act), or otherwise subject to the liabilities of that Section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended (Securities Act). The information in Item 2.02 and Exhibit 99.1 shall not be deemed incorporated by reference in any filing under the Securities Act, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description

99.1 <u>Press release dated October 8, 2020</u>
104 Cover Page Interactive Date File (embedded within the Inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

SI-BONE, INC.

Date: October 8, 2020

By:

/s/ Laura A. Francis

Laura A. Francis Chief Operating Officer and Chief Financial Officer (Principal Financial and Accounting Officer)



SI-BONE Reports Preliminary Unaudited Revenue for the Third Quarter 2020 and Provides Full Year 2020 Revenue Guidance

SANTA CLARA, Calif. October 8, 2020 - SI-BONE, Inc. (Nasdaq:SIBN), a medical device company dedicated to solving musculoskeletal disorders of the sacropelvic anatomy, today announced preliminary unaudited revenue for the three and nine months ended September 30, 2020.

Preliminary and unaudited revenue for the three months ended September 30, 2020 is expected to be in the range of \$20.1-\$20.4 million, reflecting growth of 24%-26% compared to the prior year period. U.S. revenue is expected to be in the range of \$18.7-\$18.9 million, reflecting growth of 26%-27% compared to the prior year period. International revenue is expected to be in the range of \$1.4-\$1.5 million.

Preliminary and unaudited revenue for the nine months ended September 30, 2020 is expected to be in the range of \$51.0-\$51.3 million, reflecting growth of approximately 7%-8% over the prior year period. U.S. revenue is expected to be in the range of \$47.2-\$47.4 million, reflecting growth of approximately 9% compared to the prior year period. International revenue is expected to be in the range of \$3.8-\$3.9 million.

The three and nine months ended September 30, 2020 revenue included in this release are preliminary and prior to the completion of SI-BONE's financial closing procedures and therefore may be subject to adjustment. SI-BONE expects to provide third quarter 2020 financial results during its third quarter 2020 earnings call on November 2, 2020.

On October 1, 2020, Anthem published an update to its sacroiliac joint fusion medical policy. This update retained policy language limiting indications for iFuse to cases involving a history of pelvic trauma, similar to earlier versions of the policy, and did not reflect the feedback from specialists Anthem received in mid-2020, or other substantive updates to the evidence base. Anthem has communicated to SI-BONE that this feedback and additional evidence may be taken into consideration in future policy updates in 2021.

"We are pleased with the rebound in our business during the third quarter, driven by underlying momentum in our business from a number of catalysts, including better reimbursement coverage and payment rates to surgeons, sales force hiring and investments in surgeon training," said Jeff Dunn, President, Chief Executive Officer and Chairman of SI-BONE.

2020 Revenue Guidance

SI-BONE expects full year 2020 revenue to be in the range of \$73.0-\$74.0 million, representing growth of approximately 8%-10% over 2019 revenue. SI-BONE remains cautious in the fourth quarter based upon the impact of COVID-19 on its business.

About SI-BONE, Inc.

SI-BONE is a medical device company that pioneered minimally invasive surgery of the SI joint with the iFuse Implant System. Studies have shown that the SI joint can be a source of pain in 15% to 30% of chronic low back pain. The iFuse ImplantTM, commercially available since 2009, is the only SI joint fusion device supported by multiple prospective clinical studies, including two randomized controlled trials, showing improved pain, patient function and quality of life resulting from treatment. There are over 85 peer-reviewed publications demonstrating the safety, durable effectiveness, and biomechanical and economic benefits unique to the iFuse Implant (www.sibone.com/results). This body of evidence has enabled multiple government and private insurance payors to establish coverage of the SI joint fusion procedure exclusively when performed with the iFuse Implant System.

The iFuse Implant System is intended for sacroiliac fusion for conditions including sacroiliac joint dysfunction that is a direct result of sacroiliac joint disruption and degenerative sacroiliits. This includes conditions whose symptoms began during pregnancy or in the peripartum period and have persisted postpartum for more than 6 months. The iFuse Implant System is also intended for sacroiliac fusion to augment stabilization and immobilization of the sacroiliac joint in skeletally mature patients undergoing sacropelvic fixation as part of a lumbar or thoracolumbar fusion. In addition, the iFuse Implant System is intended for sacroiliac fusion in acute, non-acute, and non-traumatic fractures involving the sacroiliac joint. There are potential risks associated with the iFuse Implant System. It may not be appropriate for all patients and all patients may not benefit.

SI-BONE and iFuse Implant System are registered trademarks of SI-BONE, Inc. ©2020 SI-BONE, Inc. All Rights Reserved. 10853-10082020

Forward Looking Statements

The preliminary unaudited financial results, and statements regarding SI-BONE's continued growth and financial outlook, contained in this press release are "forward-looking" statements. These forward-looking statements are based on SI-BONE's current expectations and inherently involve significant risks and uncertainties and, with respect to the preliminary unaudited financial results, are subject to quarterend closing adjustments. These forward-looking statements are based on SI-BONE's current expectations and inherently involve significant risks and uncertainties. These forward-looking statements are based on SI-BONE's current expectations and inherently involve significant risks and uncertainties. These risks include the impact that the COVID-19 pandemic will have on the ability and desire of patients and physicians to undergo procedures using the iFuse Implant System, the duration of the COVID-19 pandemic, whether the COVID-19 pandemic will increase in severity in the future, and SI-BONE's ability to increase demand for iFuse and obtain favorable coverage and reimbursement determinations from third-party payors. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of the risks and uncertainties, many of which are described in the company's most recent filings on Form 10-K and Form 10-Q, and the company's other filings with the Securities and Exchange Commission (SEC) available at the SEC's Internet site (<u>www.sec.gov</u>), especially under the caption "Risk Factors". SI-BONE does not undertake any obligation to update forward-looking statements and expressly disclaims any obligations or undertaking to release publicly any updates or revisions to any forward-looking statements contained herein, except as required by law.

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