



## **SI-BONE To Report Third Quarter 2019 Financial Results on November 11, 2019**

October 28, 2019

SANTA CLARA, Calif., Oct. 28, 2019 (GLOBE NEWSWIRE) -- SI-BONE, Inc. (Nasdaq: SIBN), a medical device company dedicated to solving musculoskeletal disorders of the sacropelvic anatomy, today announced it will report financial results for the third quarter of 2019 after market close on Monday, November 11, 2019. The company's management will host a corresponding conference call beginning at 1:30 p.m. Pacific Time / 4:30 p.m. Eastern Time.

Investors interested in listening to the conference call may do so by dialing (866) 470-1968 for domestic callers or (409) 217-8248 for international callers, using conference ID: 8881559. A live and archived webcast will be available on the "Investors" section of the company's website at: [www.si-bone.com](http://www.si-bone.com).

### **About SI-BONE**

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 Implant™, commercially available since 2009, is the only SI joint fusion device supported by multiple prospective clinical studies, including two RCTs, showing improved pain, patient function and quality of life resulting from treatment. There are over 70 peer-reviewed publications demonstrating the safety, durable effectiveness, and biomechanical and economic benefits unique to the iFuse Implant ([www.si-bone.com/results](http://www.si-bone.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 sacroiliitis. 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 immobilization and stabilization of the sacroiliac joint in skeletally mature patients undergoing sacropelvic fixation as part of a lumbar or thoracolumbar fusion.

There are potential risks associated with the iFuse Implant System. It may not be appropriate for all patients and all patients may not benefit. Rx Only. For information about the risks, visit: [www.si-bone.com/risks](http://www.si-bone.com/risks)

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Source: SI-BONE, Inc.