



SI-BONE To Report First Quarter 2019 Financial Results on May 8, 2019

April 24, 2019

SANTA CLARA, Calif., April 24, 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 first quarter of 2019 after market close on Wednesday, May 8, 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: 6458389. A live and archived webcast will be available on the "Investors" section of the company's website at: www.SI-BONE.com.

About SI-BONE, Inc.

SI-BONE is a medical device company that pioneered the iFuse Implant System, a minimally invasive surgical system for fusion of the sacroiliac joint to treat sacroiliac joint dysfunction. The SI joint is the last major joint with a clinically proven surgical treatment. The iFuse Implant, commercially available since 2009, is the only SI joint fusion device supported by multiple prospective clinical studies showing improved pain, patient function and quality of life resulting from treatment. There are over 65 peer-reviewed publications supporting the safety, durable effectiveness, and biomechanical and economic benefits unique to the iFuse Implant (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.

SI-BONE and *iFuse Implant System* are registered trademarks of SI-BONE, Inc. iFuse Bedrock is a trademark of SI-BONE, Inc. ©2019 SI-BONE, Inc. All Rights Reserved.

Investor Contact:

Lynn Lewis or Carrie Mendivil

investors@SI-BONE.com



Source: SI-BONE, Inc.