The Sacroiliac Joint As a Cause of Lower Back Pain



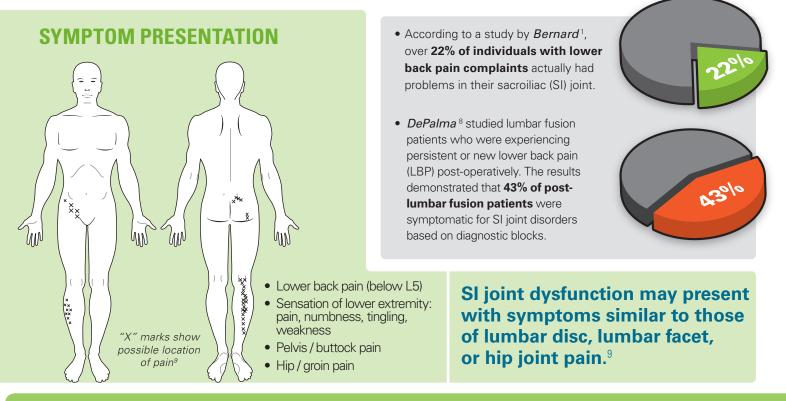
t 408.207.0700 f 408.577.8312 e info@si-bone.com www.si-bone.com

> SI-BONE | **iFuse** Implant System Minimally Invasive Sacroiliac Joint Surgery

# **SI Joint in Lower Back Pain**

15-30% of patients with lower back pain had symptoms related to the sacroiliac (SI) joint.<sup>1-5</sup> Yet, most published studies on differential diagnosis for lower back pain fail to consider the SI joint in their algorithms.<sup>6,7</sup>

**CAUSES:** • Degenerative disease • History of trauma • Pregnancy/childbirth • Prior lumbar fusion



### DIAGNOSIS

Provocative Tests<sup>10,11</sup> followed by Diagnostic Injection(s) when indicated, are recommended to confirm pain is coming from the SI joint.



#### SI JOINT DIAGNOSTIC INJECTIONS



- Fluoroscopic guidance
- 0.3 0.5 cc contrast medium
- 1.4 2.0 cc local anesthetic
- Post-injection pain reduction: ≥ 50% SIJ is likely the source of pain

< 50% Should consider other pain sources, but SIJ may be a component

\* Flexion, Abduction, External Rotation

### **NON-SURGICAL TREATMENT**

Treatment goals for SI joint pain include reducing symptoms and improving patient function. Non-surgical treatment options may include:

- Oral pain medications (NSAIDs, opioids, etc.)
- Chiropractic treatments

Therapeutic SI joint

- SI belt
- Physical therapy
- RF ablation

iniections

## iFuse Implant<sup>™</sup>

The iFuse Implant, with its patented triangular implant design, has produced **unparalleled clinical results.**<sup>12</sup> Multiple published, peer-reviewed articles demonstrate the safety and effectiveness of the iFuse Implant, including results from prospective multicenter clinical trials.<sup>13-16</sup> The iFuse Implant is the only SI joint device system with clinical studies demonstrating that treatment improved pain, patient function, and quality of life.

## **SURGERY: SI JOINT FUSION**

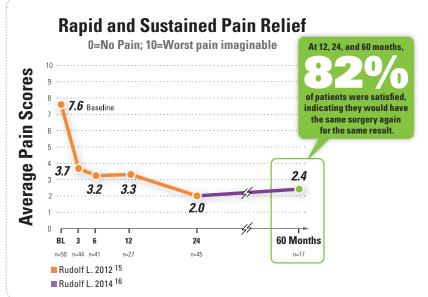
Patients refractory to non-surgical management may benefit from minimally invasive surgical (MIS) fusion of the SI joint.





#### UNIQUE TRIANGULAR DESIGN

- Triangular implant profile minimizes rotation and an interference fit minimizes micromotion
- Implant's large porous surface area allows for bony bony ongrowth/ ingrowth<sup>19</sup>
- Rigid titanium construction and implant geometry provide immediate stabilization



If you have patients who have failed non-surgical treatment, please call 866-811-4159 to find an iFuse trained surgeon.

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#### SI-BONE, Inc.

471 El Camino Real Santa Clara, CA 95050 USA

- t 408.207.0700
- f 408.557.8312
- e info@si-bone.com

#### www.si-bone.com

The iFuse Implant System<sup>®</sup> 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. There are potential risks associated with the iFuse Implant System. It may not be appropriate for all patients and all patients may not benefit. For information about the risks, visit *www.si-bone.com* Rx Only.

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