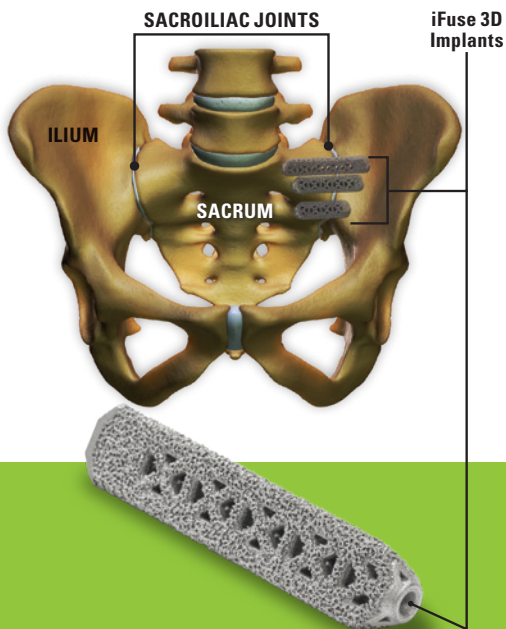


Minimally Invasive SI Joint Fusion with the iFuse 3D™ Implant System



The **iFuse 3D™ Implant System** is intended for sacroiliac (SI) joint fusion for conditions including SI joint dysfunction that is a direct result of SI joint disruption or degenerative sacroiliitis.

The procedure involves the insertion of three small, triangular, 3D-printed titanium implants across the SI joint, and is designed to stabilize and fuse the SI joint. The procedure is done through a small incision and takes approximately one hour.

Clinical studies have demonstrated that treatment with the Fuse Implant System improved pain, patient function, and quality of life.^(1,2)

1 Polly, D.W. et al. *Int J Spine Surg*. 2016 Aug 23;10:28.

2 Dengler J, et al. *J Bone Joint Surg Am*. 2019;101(5):400-11.

Since 2009, SI-BONE has been a trailblazer in the treatment of SI joint dysfunction, offering products with supporting clinical evidence for the effective management of this condition. We have developed a range of cutting-edge solutions that have evolved over the years to cater to the individual needs of patients and providers.

To learn more about SI joint dysfunction, the iFuse 3D Implant System, and other patient resources, scan the QRC below:



The **iFuse 3D™ Implant System** is intended for sacroiliac fusion for the following conditions:

- 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.
- To augment immobilization and stabilization of the sacroiliac joint in skeletally mature patients undergoing sacropelvic fixation as part of a lumbar or thoracolumbar fusion.
- Acute, non-acute, and non-traumatic fractures involving the sacroiliac joint.

If present, a pelvic fracture should be stabilized prior to the use of iFuse implants.

Healthcare professionals should refer to the Instructions For Use for indications, contraindications, warnings, and precautions at www.si-bone.com/label.

There are potential risks associated with iFuse procedures. They may not be appropriate for all patients and all patients may not benefit. For information about the risks, visit www.si-bone.com/risks.

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iFuse 3D™
Implant System

SI-BONE®

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Patients

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