Indications

The SI BONE iFuse INTRA Ti Implant System is intended for fusion of the sacroiliac joint for 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.

Healthcare professionals should refer to iFuse INTRA Ti Instructions For Use (503148), for contraindications, warnings and precautions at https://si-bone.com/label

There are potential risks associated with the iFuse INTRA Ti 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/risks.

Complaints and adverse events relating to use of the procedure and/or device should be reported to SI BONE, Inc. Toll Free: (855) 511-1545 or E mail qara@si bone.com