



Indications for use:

Bone fractures, Osteotomies, Arthrodesis, Osteochondritis, and tendon re-attachment. It is intended for but not limited to hand surgery, plastic surgery, and podiatric surgery, but is not intended for use in the spine.

BioPro Go-EZ Screw™

Contra-indications:

1. Comminuted bone surface that would mitigate against screw placement.
2. Pathologic conditions of bone such as osteopenia which would impair the ability to securely fix the screw.
3. Foreign body sensitivity to metals specifically titanium. Where material sensitivity is suspected, appropriate tests should be made prior to implantation.

Sterile:

Sterilized with ethylene oxide gas. Caution: Do not use if package is open or damaged. This is a single use device. Re-use of this device can result in the transfer of materials not limited to bone, tissue, blood, or infectious disease. The device is provided sterile and re-sterilization of the device has not been validated.

The Go-Ez Screw has been evaluated for safety and compatibility in the MR environment and is MR conditional. Contact BioPro for MR parameters. Contact surgeon if a change in performance or pain level is noticed.

For a complete glossary of symbols, please visit www.bioproimplants.com/IFU. To receive a printed version within 5 business days, please contact orders@bioproimplants.com



BP090508 05



BioPro, Inc.
2929 Lapeer Road
Port Huron, MI 48060
810-982-7777 (P)
810-982-7794 (F)



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