

Coracoid Bone Conserving Acromioclavicular Joint Reconstruction

Using ToggleLoc™ Device with ZipLoop™ Technology

Surgical Technique
by Peter J. Evans, MD, PhD

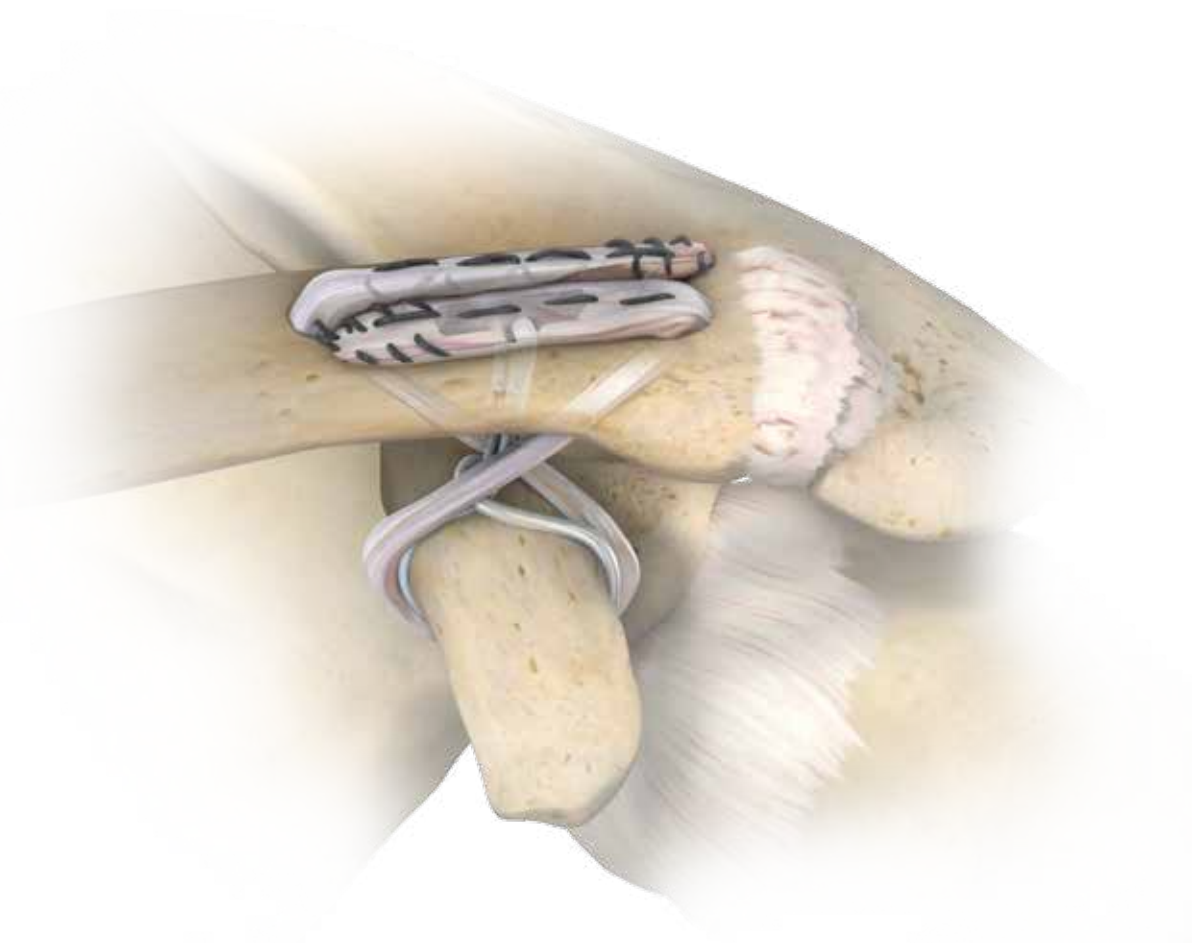


Table of Contents

- Surgical Technique Indications 2
- Acute vs. Chronic Reconstruction 2
- Patient Positioning and Dissection 2
- Pass the Implant 3
- Create the Slip Knot 4
- Prepare the Clavicle 5
- Reduce the AC Joint..... 7
- Chronic Dislocation of the ACJ–Passing the Graft..... 7
- Graft Placement 8
- Closure..... 9
- Post-operative Protocol 9
- Ordering Information 10
- Indications For Use 11
- Contraindications 11

Surgical Technique



Figure 1



Figure 2

Surgical Technique Indications

This technique is intended for treatment of acute or chronic acromioclavicular joint (ACJ) dislocation.

Acute vs. Chronic Reconstruction

When treated within four weeks of dislocation this technique can be used with the ToggleLoc with ZipLoop Technology alone, but for more chronic conditions, a tendon graft used in conjunction is warranted. In very chronic cases the distal clavicle is often resected due to degenerative changes in the ACJ.

Patient Positioning and Dissection

Position the patient in a beach chair position under general anesthesia so that fluoroscopy can be positioned to confirm ACJ reduction (Figure 1). A supraclavicular anaesthetic block can aid with postoperative pain control. A 2.5 cm oblique incision is made spanning the clavicle just superior to the coracoid in Langer's lines, which allows for cosmesis (Figure 2). A slightly longer incision is required if the distal clavicle requires excision. Alternatively, a transverse incision can be made along the inferior border of the clavicle.



Figure 3

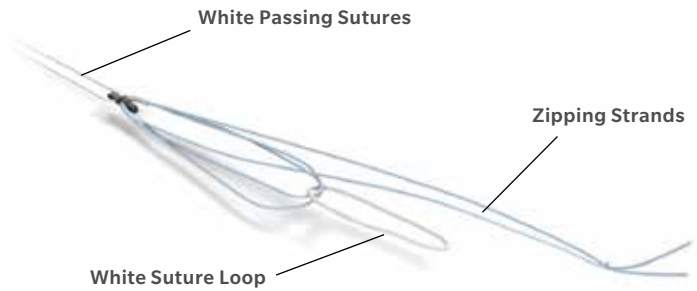


Figure 4a

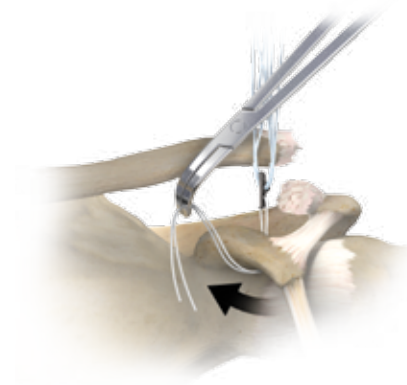


Figure 4

Patient Positioning and Dissection (cont.)

Subcutaneous sensory nerve branches are identified and protected. Elevate the skin flaps and incise the deltotrachezial fascia. Elevate the deltoid off of the clavicle to identify the subdeltoid space (Figure 3). Blunt dissection is used to identify the base of the coracoid process. Incise the fascia on the lateral and medial aspect of the coracoid.

Pass the Implant

Deliver a hemostat or other blunt curved instrument underneath the inferior aspect of the coracoid process from medial to lateral. One should feel the transition to the neck of the glenoid from the anterior scapula to assure placement at the base of the coracoid. With the curved instrument, attach the white passing sutures from the ToggleLoc device with ZipLoop Technology with continuous loop and shuttle them medially (Figure 4 & 4a).

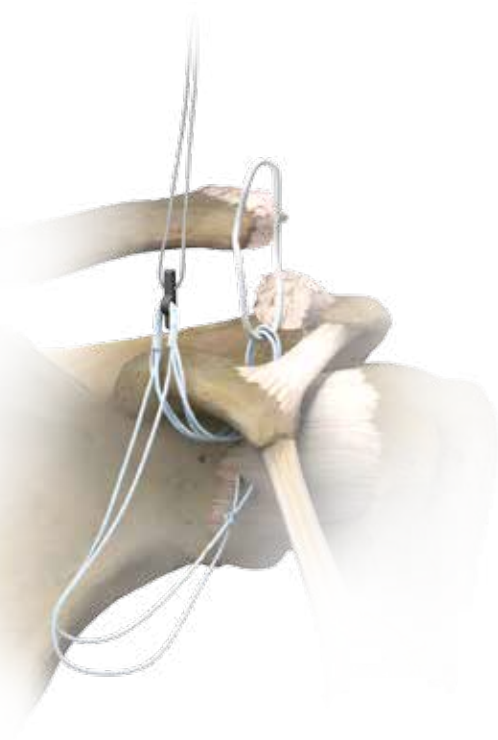


Figure 5

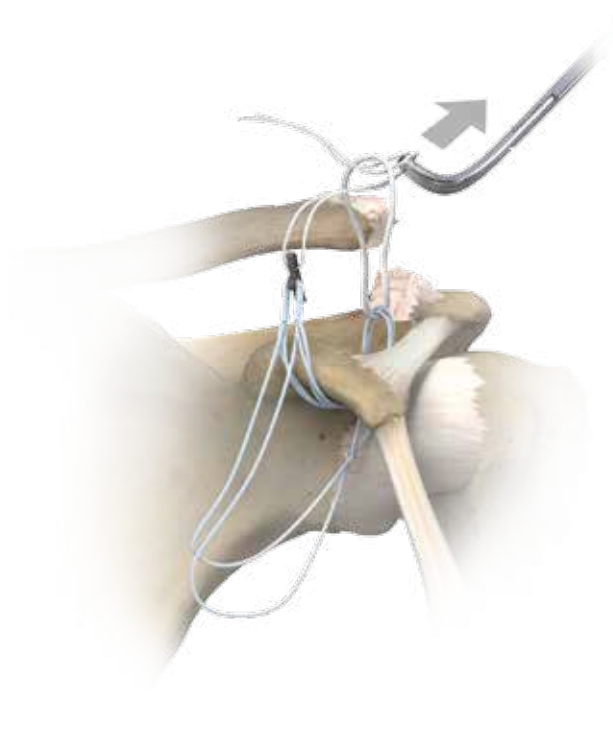


Figure 6

Pass the Implant (cont.)

Ensure the white suture loop remains on the lateral side and the MaxBraid™ zipping strands exit medially (Figure 5).

Create the Slip Knot

Next, deliver a hemostat through the solid white suture loop on the lateral side and capture the white passing sutures from the ToggleLoc device with ZipLoop Technology from the medial side. Pull the white lead sutures through the white suture loop to create the slip knot (a.k.a “luggage tag”) around the coracoid (Figure 6).

ⓘ **Note:** Creating the luggage tag completely eliminates the need for coracoid drilling.



Figure 7

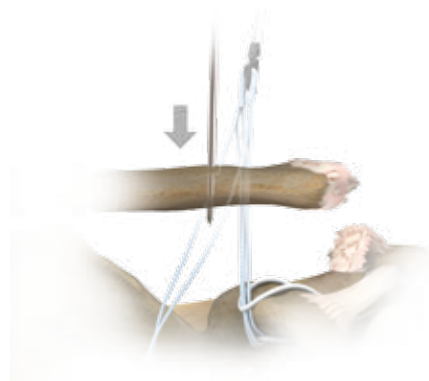


Figure 8

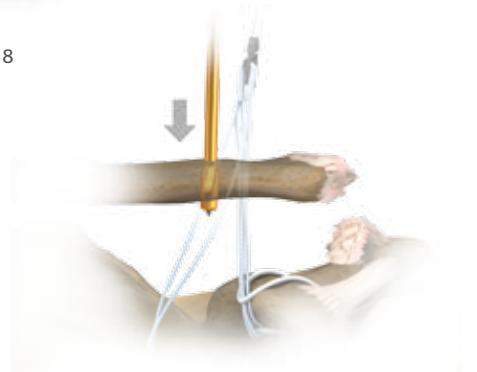


Figure 9

Create the Slip Knot (cont.)

Make sure to pass the lead sutures, ToggleLoc button, and zipping strands through the solid white suture loop and assure the solid white loop is positioned under the coracoid as its larger diameter will distribute forces optimally (Figure 7). Leave the hemostat clipped to the white passing suture as preparation of the clavicle takes place.

Prepare the Clavicle

Drill a 2.4 mm guide pin bicortically into the clavicle directly superior to the coracoid. Place a 4.5 mm cannulated reamer over the guide pin and drill bicortically (Figures 8 & 9). Leaving the 4.5 mm reamer in place, pass a Nitinol loop (or any monofilament suture) through the cannulation of the 4.5 mm reamer. This will be utilized to shuttle the leading passing strands of the ToggleLoc to the top of the clavicle.

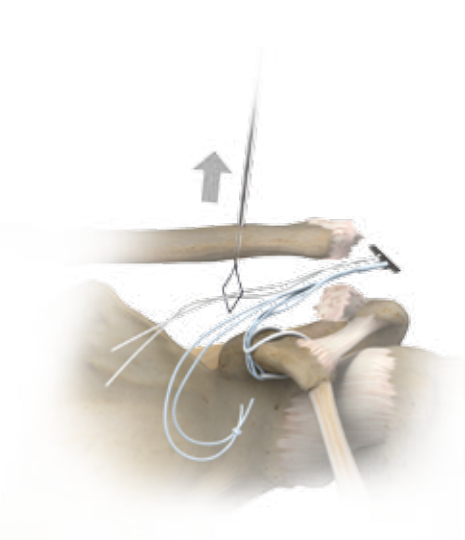


Figure 10



Figure 10a

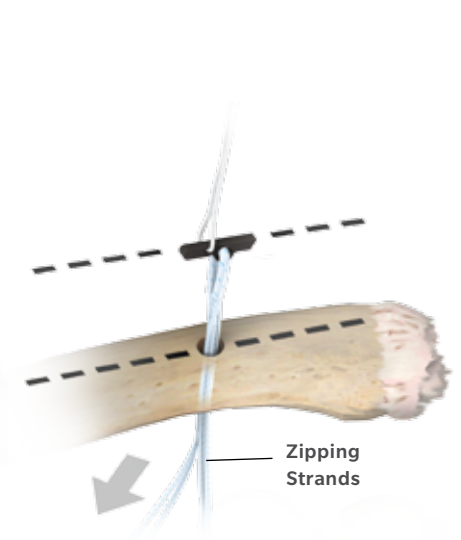


Figure 11

Prepare the Clavicle (cont.)

Release the white passing sutures from the hemostat. Utilize the Nitinol wire to shuttle the leading white passing strand of the ToggleLoc device with ZipLoop Technology and pass them from inferiorly through the clavicle and out superiorly, while simultaneously pulling the ToggleLoc button through the clavicle (Figures 10 & 10a).

The ToggleLoc button should be positioned collinear to the long axis of the clavicle (Figure 11). The zipping strands of the ZipLoop must remain infraclavicular.

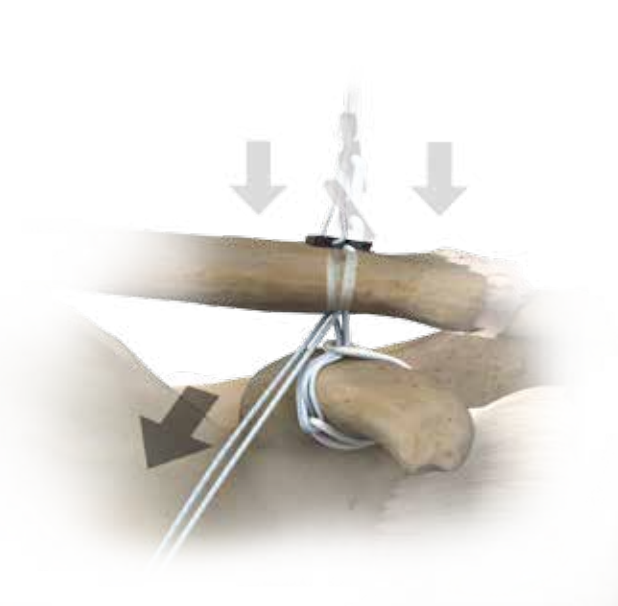


Figure 12

Reduce the AC Joint

Pull the ToggleLoc button until flush with the bone. Clavicle reduction is now performed by applying downward pressure on the clavicle and upward pressure on the elbow. The ACJ is over-reduced by 5 mm. The zipping strands of the ToggleLoc with ZipLoop are now pulled using the ZipLoop puller to achieve equal tension on both strands, taking up slack until desired reduction is achieved (Figure 12).



Figure 13

Chronic Dislocation of the ACJ—Passing the Graft

In the chronic case, a tendon graft should be passed around the coracoid. The graft can be an autograft or allograft based upon surgeon preference (Figure 13). This can be done simultaneously while the white passing sutures of the ToggleLoc with ZipLoop are shuttled from lateral to medial, as described earlier in the technique.

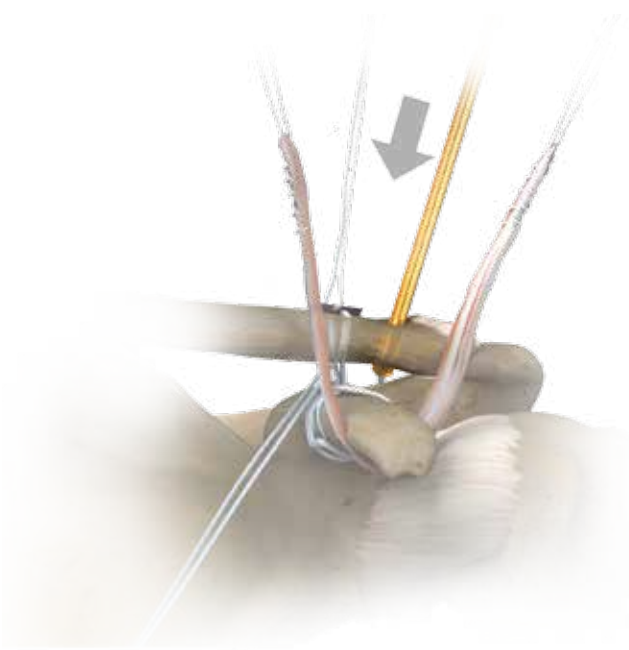


Figure 14

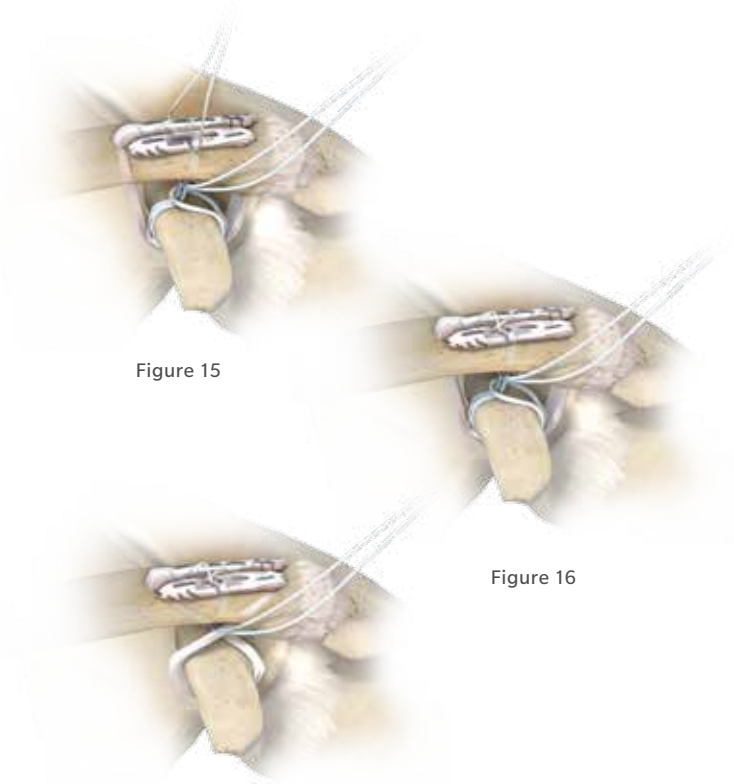


Figure 15

Figure 16

Figure 16a

Graft Placement

Two additional 3.2 mm drill holes are required to place the graft. One drill hole should be placed medial to the ToggleLoc with ZipLoop hole and the second should be placed lateral to the ToggleLoc with ZipLoop hole. Drill a 2.4 mm guide pin into the clavicle bicortically. Then drill over the pin with a 3.2 mm cannulated reamer (Figure 14).

Note: Each drill hole should be separated by 1.0-1.5 cm, recreating the anatomic locations of the coracoclavicular ligaments. Each limb of the tendon graft is passed from inferior to superior through the clavicle. A Nitinol wire can be utilized to shuttle graft suture limbs to the superior clavicle.

Once each limb of the graft is passed, the clavicle is reduced and the ZipLoop is tightened. The graft ends are then crossed parallel to each other over the ToggleLoc button. The ends should be maximally tensioned and sewn together upon exiting the clavicle with a free #2 MaxBraid suture in an interrupted figure-eight fashion (Figure 15).

The MaxBraid passing suture from the ToggleLoc implant can be tied over the tendon graft to secure the two limbs for additional fixation (Figure 16).

Optionally, crossing the graft limbs above the coracoid before entering the clavicle can create a more anatomic trajectory (Figure 16a). In reconstructive cases where not enough distal clavicle remains, two holes can be used, placing the ZipLoop and graft together through one of the holes.

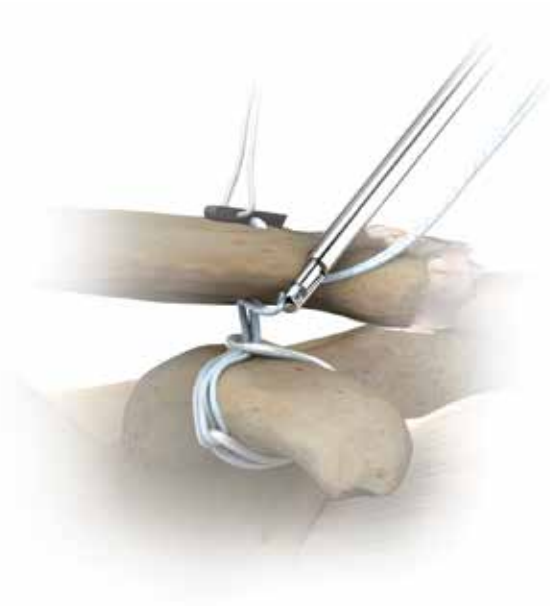


Figure 17

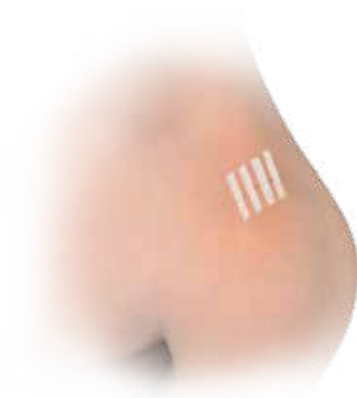


Figure 18



Figure 19: Pre-Op

Figure 19a: Post-Op

Closure

Insert the pull sutures into the Super MaxBraid cutter and advance the cutter to the inferior surface of the clavicle. Compress the trigger of the cutter until both strands of the sutures are severed (Figure 17). A layered closure is performed with absorbable sutures and a subcuticular skin closure with steri-strips for an optimal cosmetic result (Figure 18). The procedure is complete (Figures 19 & 19a).

Post-operative Protocol

A shoulder immobilizer is utilized for a period of six weeks, and waist level activities such as eating or clerical work are allowed. Range of motion is restored from weeks six to twelve. Progressive resistance exercises are initiated after twelve weeks. Overhead sport and impact loading activities are allowed after six months.

Ordering Information

Implant

Part Number	Size	Description
904756	–	ToggleLoc Device with ZipLoop Technology with Continuous Loop

Instrumentation

Part Number	Size	Description
904837	–	AC Joint Disposable Kit
904776	–	ZipLoop Puller
904011	–	Sterile Nitinol Loop-10pk
904004	2.3 mm	SpeedPass™ Suture Retriever Straight, Disposable
948084	3.2 mm	Cannulated Drill

INDICATIONS FOR USE

The ToggleLoc System devices, except the ToggleLoc XL device, are intended for soft tissue to bone fixation for the following indications:

Shoulder

Bankart lesion repair
 SLAP lesion repairs
 Acromio-clavicular repair
 Capsular shift/capsulolabral reconstruction
 Deltoid repair
 Rotator cuff tear repair
 Biceps Tenodesis

Foot and Ankle

Medial/lateral repair and reconstruction
 Mid- and forefoot repair
 Hallux valgus reconstruction
 Metatarsal ligament/tendon repair or reconstruction
 Achilles tendon repair
 Ankle Syndesmosis fixation (Syndesmosis disruptions) and as an adjunct in connection with trauma hardware for Weber B and C ankle fractures (only for ToggleLoc with Tophat/ZipTight Fixation Devices)

Elbow

Ulnar or radial collateral ligament reconstruction
 Lateral epicondylitis repair
 Biceps tendon reattachment

Knee

ACL/PCL repair / reconstruction
 ACL/PCL patellar bone-tendon-bone grafts
 Double-Tunnel ACL reconstruction
 Extracapsular repair: MCL, LCL, and posterior oblique ligament
 Iliotibial band tenodesis
 Patellar tendon repair
 VMO advancement
 Joint capsule closure

Hand and Wrist

Collateral ligament repair
 Scapholunate ligament reconstruction
 Tendon transfers in phalanx
 Volar plate reconstruction
 The ToggleLoc XL device is used for fixation of tendons and ligaments in cases of unanticipated intraoperative complications such as cortical breaching during orthopedic reconstruction procedures, such as Anterior Cruciate (ACL) or Posterior Cruciate (PCL) Reconstruction.

CONTRAINDICATIONS

1. Infection.
2. Patient conditions including blood supply limitations, and insufficient quantity or quality of bone or soft tissue.
3. Patients with mental or neurologic conditions who are unwilling or incapable of following postoperative care instructions.
4. Foreign body sensitivity. Where material sensitivity is suspected, testing is to be completed prior to implantation of the device.

This material is intended for health care professionals and the Zimmer Biomet sales force only. Distribution to any other recipient is prohibited. All content herein is protected by copyright, trademarks and other intellectual property rights owned by or licensed to Zimmer Biomet or its affiliates unless otherwise indicated. This material must not be redistributed, duplicated or disclosed, in whole or in part, without the express written consent of Zimmer Biomet.

Check for country product clearances and reference product specific instructions for use. For complete product information, including indications, contraindications, warnings, precautions, and potential adverse effects, see the package insert and Zimmer Biomet's website.


This technique was prepared in conjunction with a licensed health care professional. Zimmer Biomet does not practice medicine and does not recommend any particular orthopedic implant or surgical technique for use on a specific patient. The surgeon is responsible for determining the appropriate device(s) and technique(s) for each individual patient.

Not for distribution in France.

©2016 Zimmer Biomet

 **Authorized Representative**
Biomet UK Ltd.
Waterton Industrial Estate
Bridgend, South Wales
CF31 3XA
UK

 **ZIMMER BIOMET**
Your progress. Our promise.™

 **Legal Manufacturer**
Biomet Sports Medicine
P.O. Box 587
56 E. Bell Drive
Warsaw, Indiana 46581
USA

www.zimmerbiomet.com

 0086

CE mark on a surgical technique is not valid unless there is a CE mark on the product label.

0401.1-GLBL-en-REV0416