Surgical Technique

Modified Garden Procedure:
Lateral Epicondylitis Repair

Surgical Protocol by
Jeffrey Nacht, M.D.
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Introduction
For the treatment of recalcitrant lateral epicondylitis refractory to non-operative measures, multiple operative techniques have been proposed. Most procedures involve debridement of the damaged segment of the common extensor origin and some type of repair of the remaining functional tendon. The technique outlined in this guide follows the principles of the Garden Procedure first proposed by R. S. Garden in 1961 in a modified and simplified approach. Utilizing the efficiency, ease of deployment, and secure fixation of the JuggerKnot anchor allows a straightforward surgical procedure to accomplish these steps.

In patients with smaller elbows, or if the preference is to avoid complete detachment of the extensor origin, the technique by Nirschl and modified by Plancher and Bishai may be substituted, which is noted on page 8.

Preoperative Preparation and Positioning
The patient is positioned supine with a “Hand Table” attachment to the operating table. A layer of bath blankets is placed beneath the elbow and forearm to position the arm parallel to the table with the elbow flexed 70 to 90 degrees (Figure 1).

A tourniquet is applied high on the operative arm. The limb is exsanguinated and the incision site injected with local anesthetic.

This material represents the surgical technique utilized by Jeffrey Nacht, M.D. Biomet does not practice medicine. The treating surgeon is responsible for determining the appropriate treatment, technique(s), and product(s) for each individual patient.
Incision
The incision is a 6 cm gently curving posterolateral approach beginning 2 cm proximal and 1 cm posterior to the lateral epicondylar prominence and following the skin fold distally. The incision is carried 1 cm posterior to the epicondyle and extended into the forearm for 3 cm, along the posterior edge of the radiocapitellar joint (Figure 2).

Dissection
Subcutaneous dissection is carried down to the deep antibrachial fascial layer, which is marked for later closure (Figure 3). Retracting this layer exposes the common origin of the Extensor Carpi Ulnaris, Extensor Carpi Radialis Brevis and Anconeus (Figure 4). The edge of this origin is divided and elevated with a needle tipped cautery (Figure 5).
Dissection (cont.)

This creates a curved, tongue-shaped full thickness layer, which can be retracted distally. Dissection is carried down to the superior posterior edge of the epicondyle, which invariably uncovers a large hypertrophic spur of the lateral humeral epicondyle (Figure 6). The full thickness flap is dissected distally off the bone to the level of the joint capsule. Although the joint is not intentionally opened, sometimes the capsule of the joint will be violated. If the joint was accidentally opened or found to be opened, it will seal with a proper closure using nonabsorbable sutures.¹ A stay suture is placed in the flap and clamped distally to the drapes to prepare the epicondyle (Figure 6a).

Bone Preparation

Next a small oscillating saw or rongeur is used to debride the hypertrophic bone and to create a flat surface for reattachment of the tendon (Figure 7). The deep surface of the tendon is inspected for degenerative angiofibrotic tendinosis tissue, which is dissected away to expose healthy deep tendon surface.

The tendon will be placed back on the bony surface and its planned reattachment site is marked with a marker or cautery for anchor placement (Figure 8). This avoids over-tensioning the tendon as the sutures are tied. The flap is again retracted to view the marked location for the two JuggerKnot anchors that will be deployed into the epicondyle in a line perpendicular to the long axis of the tendon (Figure 8a).
Placement of the JuggerKnot Guide

Place two JuggerKnot Soft Anchors-1.4mm Short. The first anchor should be placed slightly more anteriorly and the second anchor should be placed more posteriorly into the prepared bed of bone. Start by placing the JuggerKnot guide onto the bone surface (Figure 9).

Without moving the guide insert the JuggerKnot 1.4mm Short drill bit into the power drill to the proximal laser-etch line to ensure appropriate depth as the collar of the drill contacts the back of the guide. Advance the drill until contact is made with the guide (Figures 10, 10a).
Insert Anchor

Remove the drill. Caution must be taken to maintain precise guide position over the pilot hole during removal. While maintaining the guide position firmly against the bone, align the luer cap on the implant inserter handle with the slot of the guide, and insert the JugglerKnot Soft Anchor through the guide and into the pilot hole. Lightly mallet to fully seat the anchor into bone (Figures 11, 11a, 11b).

Deploy Anchor

Once the anchor has been fully seated into the reattachment site on the surface of the epicondyle, lightly pull back on the anchor inserter handle to set the anchor (Figures 12, 12a).
**Deploy Anchor (cont.)**

Release the suture from the handle by unscrewing the suture retention feature and remove the needles from the middle of the guide (Figure 13). Remove the anchor inserter and guide. Lightly pull on both sutures to set the anchor and verify the sutures slide smoothly.

Follow the same steps for the deployment of the second anchor (Figure 14).

**Attachment of Tendon**

Using the needles attached to the JuggerKnot Soft Anchor (2,T-5 Needles), the MaxBraid sutures are passed from the deep to the superficial surface of the tendon and tied (Figure 15). **The sutures are NOT CUT.** Next, the same sutures are passed into the proximal edge of the common extensor origin, tucking this layer over the reattached tendon edge in a “pants-over-vest” manner (Figure 16). This will cover the bone left exposed by the distally displaced tendon origin.
Attachment of Tendon (cont.)

Next, the common extensor origin repair is reinforced and completed with a running suture of 2-Vicryl or similar absorbable suture, following the curve of the detachment site. The antebrachial fascial layer is then repaired with 2-0 Vicryl (Figure 17).

Closure

The deep antebrachial fascia is repaired with 2-0 Vicryl (Figure 18). The subcutaneous and skin closure is completed using the surgeon’s preferred technique (Figure 19). A light padded dressing is applied over the olecranon and lateral elbow, followed by a layer of sterile Webril. A posterior plaster splint is applied and secured with an elastic wrap. The elbow is positioned at 90 degrees of flexion in a sling.

Postoperative Care

The dressing and splint are removed at one week postoperatively. This repair is very secure and allows for early mobilization. Hand use is encouraged immediately postoperatively. After one week, a light dressing is applied and early elbow motion is encouraged. At three weeks the patient may begin physical and hand therapy to regain range of motion and dexterity. At four weeks strengthening may begin.
Alternate Technique

As noted in the introduction, in patients with smaller elbows, or if the preference is to avoid complete detachment of the extensor origin, the technique by Nirschl and modified by Plancher and Bishai may be substituted. Their technique involves an identical approach down to the common extensor origin. Instead of creating a tongue-shaped flap, the origin is split longitudinally between the ECRB/ECRL and the EDC along their fibers, at its central point and retracted to expose the underlying epicondylar osteophyte (Figure 20). The epicondyle is similarly debrided to a flat surface through the interval in the tendon layer and all necrotic angiofibroblastic tissue and torn tendon fibers are excised along with any granulation tissue (Figure 21).

Next two JuggerKnot anchors are deployed into the bony surface of the epicondyle, one under each edge of the tendon (Figure 22). The sutures are passed through the two limbs of the extensor origin and left untied while the longitudinal split in the tendon is repaired with a running suture of braided polyester (non-absorbable) or Vicryl (absorbable) size 0 suture, according to the preference of the surgeon. The sutures from the anchors are then tied over the surface of the tendon (Figure 23). The closure of the antebrachial fascia and skin are carried out as noted on page 7 (Figures 18, 19).
Ordering Information

**JuggerKnot Soft Anchor Short**

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<tr>
<th>Part Number</th>
<th>Size</th>
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<td>JuggerKnot Soft Anchor Short w/Needles Single Loaded</td>
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<td>JuggerKnot Soft Anchor Short Drill Bit (Disposable) Sterile</td>
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<td>912075</td>
<td>1.4 mm Short</td>
<td>JuggerKnot Soft Anchor Short Implant System</td>
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References

Indications for use and contraindications for use for the JuggerKnot Soft Anchor–1.4 mm Short are as follows:

INDICATIONS
The JuggerKnot Soft Anchors are intended for soft tissue to bone fixation for the following indications:

Shoulder
- Bankart lesion repair
- SLAP lesion repair
- 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

Elbow
- Ulnar Radial collateral ligament reconstruction
- Lateral epicondylitis repair
- Biceps tendon reattachment

Knee
- Extra-capsular 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

Hip
- Acetabular labral repair

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 or patients who are otherwise unwilling or incapable of doing so.
4. Foreign body sensitivity. Where material sensitivity is suspected, testing is to be completed prior to implantation of the device.

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