Lateral Ankle Ligamentous Instability

Surgical Technique by Jeffrey Nacht, M.D.
It’s small. It’s strong. And it's all suture.

The JuggerKnot Soft Anchor represents the next generation of suture anchor technology. The 1.4mm deployable anchor design is a completely suture-based system, and is the first of its kind.

Suture Configuration
- Loaded with #1 MaxBraid Suture—leaves a lower knot profile vs. a #2 suture

Needles
- Tapered #5 needles can be used to tie down ligaments
Minimal Size
- Smaller drill guide is designed to be less invasive to surrounding tissue
- Smaller anchor diameter allows multiple anchors to be placed
- Reduces likelihood of intersecting anchors when placing multiple anchors

Reduced Bone Removal
- The volume of bone that is removed with a 3.0mm drill is equivalent to four JuggerKnot device drill holes

Soft Material
- Soft anchor deployment system—completely suture based implant
- Implant made from #5 polyester suture
- Eliminates the possibility of rigid material loose bodies in the joint
Introduction
Several procedures have been described using the Biomet Sports Medicine JuggerKnot Soft Anchor, designed to take full advantage of its ease of use, rapid deployment, small footprint within the bone, and its ability to provide fixation in multiple layers which can save time and decrease the need for surgical exposure and dissection compared with standard techniques.

Pre-Operative Preparation and Positioning
A popliteal block is placed for postoperative analgesia before beginning the procedure. The patient is then anesthetized with regional or general technique and placed in a full lateral or semi-lateral position (surgeon’s preference) and secured with a vac-pack or similar device. A semi-lateral position is preferred to allow the addition of a diagnostic ankle arthroscopic procedure, which is included with most cases, due to the high incidence of accompanying intra-articular pathology.

Surgical Technique
A tourniquet may be placed at either the mid-calf or the mid-thigh. Generally, the mid-thigh is preferred so that the surgeon may choose to augment the repair with a local transfer of a peroneal tendon if the local tissues are found to be inadequate to accomplish the procedure. A small roll of sterile towel is placed beneath the ankle to support the inverted ankle position during the procedure.

Incision
The incision is placed as a gently curving line running approximately 1 cm anterior to the distal fibula and following its line inferiorly toward the peroneal tendons (Figure 1). Alternatively, a posteriorly placed curving incision along the posterior edge of the fibula and then curving onto the lateral border of the foot can also be used. This allows easier access to the peroneal tendons. The skin and subcutaneous tissue is divided and retracted with skin hooks or senn rakes.
Dissection

Next the inferior extensor retinaculum is dissected from the subcutaneous layer distally and slightly proximally, to prepare for later reefing. A marking pen can be used at the incision in the retinaculum to aid in finding this edge for later closure (Figure 2). After incising this layer along the marked line, it is retracted to expose the ankle capsule.

Tip: To ensure that the ankle is opened and the subtalar joint is not inadvertently violated, palpate the edge of the fibular malleolus and the ankle joint with a 22 gauge needle before making an incision.

The capsule is then incised along the anterior edge of the fibula, following the bone edge inferiorly. Before reaching the peroneal tendons, insert the blunt end of a senn rake or ragnel retractor to protect the peroneals as the incision is carried posteriorly. This step also exposes the Calcaneo-fibular ligament (CFL) which lies just beneath the peroneal tendons in this location and is easily located by "toeing-in" the senn rake (Figure 3).

The CFL is released off the fibula and the ankle is inspected for any loose bodies and debris.

By sharp dissection, the fibular periosteum is elevated off the edge of the fibula and carefully preserved for later use in the repair. The distal and anterior edge of the fibula is prepared with a 3–4 mm burr, creating a shallow decorticated trough into which the ligaments will be re-attached (Figure 4).
Placement of the JuggerKnot Short Guide
Next, place two JuggerKnot Soft Anchors. The first anchor should be placed onto the fibula where the anterior talo-fibular ligament (ATFL) should be reattached. The second anchor should be placed at the inferior tip of the fibula where the Calcaneofibular ligament (CFL) will be re-attached. Start by placing the JuggerKnot guide onto the fibula where the ATFL should be reattached (Figure 5).

Drill the Pilot Hole
Without moving the guide insert the JuggerKnot 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 6 & 6a).
**Insert the Anchor**

Remove the drill. **Note: Caution must be taken to maintain precise guide position over the pilot hole during removal.** While maintaining the guide position firmly against the bone, insert the JuggerKnot Soft Anchor through the guide and into the pilot hole. Lightly mallet to fully seat the anchor into bone (Figures 7 & 7a).
Deploy the Anchor

Once the anchor has been fully seated into the fibula, lightly pull back on the anchor inserter handle to set the anchor (Figure 9). Release the suture from the handle by unscrewing the suture retention feature and remove the needles from the middle of the guide (Figure 9a). Pull the anchor inserter handle directly back from the guide. Lightly pull on both sutures to set the anchor and verify the sutures slide (Figure 10).

Repeat these steps to place the second JuggerKnot Soft Anchor on the fibula at the location of the CFL (Figure 11).
Repair the Tendons

Use the needles attached to the JuggerKnot Soft Anchor (#5 Taper Needle) to pass the Maxbraid sutures. The anterior sutures are passed in a modified Bunnell fashion through the thickened capsule at the site of the ATFL, passing the needle twice into the ligament for each suture limb. Next, the posterior sutures are passed into the edge of the CFL in a similar manner (Figure 11a). The towel roll is now removed from beneath the ankle and the foot placed in a neutral position. While the assistant holds the ankle in a gently everted position, the sutures are tied, bringing the ligaments snugly up against the prepared bone edge at the proper tension. This completes the first layer of repair. The sutures are **NOT cut**.

The same sutures are then passed through the periosteal layer on the fibula and tied over this second layer, tucking it over the primary repair as reinforcement. **Again, the sutures are NOT cut.** (Figures 11b & 11c)
Surgical Technique

The distal edge of the Inferior Extensor Retinaculum is now pulled up over the repair and the sutures are passed through this layer and tied again, providing the “Gould” reinforcement (Figure 12). Finally, the proximal edge of the Inferior Extensor Retinaculum is drawn over top of the distal edge in a “pants-over-vest” manner to further reinforce the repair (Figure 13). The JuggerKnot sutures are cut after this last repair (Figure 14).

Testing the Repair
The repair is now evaluated by lifting the limb off the operating table. The ankle should now have enough lateral stability to resist inversion from the force gravity. Further, full ankle dorsiflexion and plantarflexion should be permitted to passive range of motion.
Closure
A running suture of 2-0 Vicryl or similar absorbable suture is then placed from anterior to posterior along the retinacular repair line. The subcutaneous and skin layers are re-approximated with the surgeon's preferred technique (Figure 15).

A well-padded posterior splint, with medial, lateral, and posterior plaster slabs secured with an elastic bandage, is prepared with an ABD pad over the heel. The ankle is held in a gently everted and dorsiflexed position as the splints set up.
**Ordering Information**

<table>
<thead>
<tr>
<th>Item Description</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>JuggerKnot Soft Anchor Short w/Needles</td>
<td>912068</td>
<td>1.4mm Single Loaded</td>
</tr>
<tr>
<td></td>
<td>912069</td>
<td>1.4mm (Package of 10)</td>
</tr>
<tr>
<td>JuggerKnot Soft Anchor Short 1.4mm Drill Bit (Disposable)</td>
<td>912071</td>
<td>Sterile</td>
</tr>
<tr>
<td>JuggerKnot Soft Anchor Short 1.4mm Guide (Reusable)</td>
<td>912072</td>
<td>Non-Sterile</td>
</tr>
</tbody>
</table>

**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 or 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.

This material is intended for health care professionals and the 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 Biomet Inc. 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 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 Biomet’s website.

This technique was prepared in conjunction with a licensed health care professional. 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.

Vicryl is a registered trademark of Johnson & Johnson Corp.