

All-Inside Meniscus Repair

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Abstract

Meniscal surgery has undergone a considerable shift in goals over the last century. While early meniscal surgery consisted of mostly total meniscectomy, recognition of the importance of this structure resulted in a shift to partial meniscectomy, and then to repair in appropriate patients. The over-reaching goal is now the preservation of meniscal tissue to minimize the risk of osteoarthritis, particularly in the young athlete. Technologic advances in arthroscopy and instrumentation have allowed the development of minimally invasive techniques, which decrease the risks associated with open surgery. While no meniscal repair technique has been demonstrated to be superior in its outcomes, the all-inside technique requires no accessory incisions and minimizes the risk to posterior structures. While the early all-inside implants have been shown to risk chondral damage, the literature demonstrates that newer suture-based implants do not share these complications, and result in the healing of appropriate tears.

Key Words: All-inside, Meniscal tear, Meniscal repair, Chondral injury

Introduction

The first open meniscal repair was performed in 1885 [1], though resection has been more common. With the advent of arthroscopy, minimally invasive techniques replaced open repairs, and provided better access to difficult areas while minimizing surgical risks. The inside-out suture repair was initially the described, and continues to be used with excellent results [2]. The outside-in repair was developed later to decrease risk of injury to posterior neurovascular structures [3]. In recent years, advances in instrument and implant technology have allowed the development of all-inside repair techniques. These rely on specialized implants, but avoid additional incisions, decreasing risk to posterior neurovascular elements, and reducing

surgical times [4]. The purpose of this review is to examine the evolution of the all-inside meniscal repair technique, with outcomes and complications.

Meniscus Anatomy, Function, And Healing:

The menisci are crescent shaped fibrocartilaginous structures situated in both the medial and lateral compartments of the knee, between the femur and tibia. Each meniscus has an anterior and posterior horn, and is attached to the tibia by the anterior and posterior meniscal roots and to the peripheral capsule by the coronary ligaments. They are triangular in cross section, conforming to both the distal femur and proximal tibia. This conformity effectively deepens the articular surfaces of the knee, providing shock absorption and contributing to stability, particularly with injury to stabilizing ligaments. This also increases the surface area for load distribution to the articular cartilage, decreasing contact stresses by converting vertical compression stresses to radially oriented hoop stresses [5, 6]. By maintaining space in the joint, the meniscus improves diffusion of synovial fluid, and provides nutrition and lubrication to the

cartilage.

The healing capacity of the meniscus is determined primarily by blood supply, as it is largely avascular and does not typically heal spontaneously. The meniscus is divided into zones in accordance with blood supply and healing capacity. The peripheral third (within 3 mm of the meniscosynovial junction) is well vascularized, as the blood supply enters the here [7]. This zone is referred to as red-red, and is mostly likely to heal. The inner third (over 5 mm from the meniscosynovial junction) receives no vascular supply, is called the white-white zone, and is least likely to heal. The middle zone, called red-white (between 3 to 5 mm from the meniscosynovial junction), has some vascularity [7, 8]. Red-red zone tears are commonly repaired in appropriate patients, while repairs of red-white zone tears are less likely to heal.

Meniscal tear characteristics also influence healing potential. Longitudinal vertical tears (including bucket handle tears and meniscocapsular tears) have the capacity to heal with repair, while degenerative or complex (multi-planar), radial, horizontal, or flap tears are much less likely. Larger and less stable meniscal tears have higher failure rates, as have those repaired more than 8 weeks from injury [9]. Lateral compartment tears are also more likely to heal than those occurring medially [9]. This may be due to increased blood supply to the posterior horn

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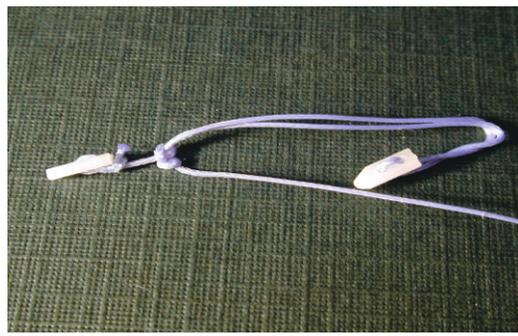


Figure 1: The FasT-Fix device has two 5 mm anchors connected by a self-locking knot of either No. 0 braided polyester or No. 0 UltraBraid suture. (copyright by author)



Figure 2: The Fast-Fix 360 has two "arrow" shaped PEEK anchors connected by No. 2-0 UltraBraid with a sliding locking knot. (copyright by author)

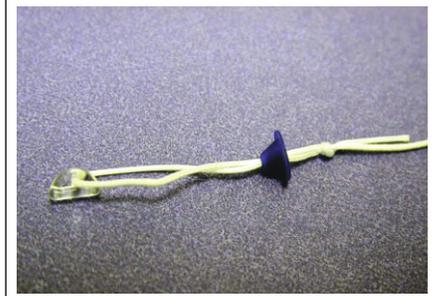


Figure 3: The RapidLoc is an adjustable suture-based device consisting of a PLLA "backstop" and either a PLLA or PDS "top hat", connected by either a No. 2-0 absorbable Panacryl or non-absorbable braided polyester suture. (copyright by author)

of the lateral meniscus.

Meniscal Repair Technique

To overcome the inherent physiologic challenges of meniscal repair, the environment and technique must be optimized. Factors controlled by the surgeon include tissue preparation, the stability of fixation, knee stability and leg alignment, and post-operative rehabilitation. Preparation should include rasping of the tear, and the perimeniscal synovium. This stimulates the healing response [10], and can allow healing of isolated stable tears without fixation, particularly with concomitant ACL reconstruction [10,11]. Some advocate trephination to create vascular access channels, which may contribute to fibrovascular healing of avascular areas [12]. The addition of fibrin clot [13, 14], platelet-rich fibrin matrix [15], and collagen matrix with bone marrow [16] have been demonstrated to aid healing. Meniscal repair with associated ACL reconstruction improves healing, possibly by increasing blood in the joint, while lack of ACL function practically assures failure from stresses on the repair [17, 18]. ACL deficiency increased the failure rates of meniscal repair from 5% to 46% [19], demonstrating the importance of stability. Normal knee alignment also is required for successful meniscus repair outcomes. Forces within the knee, and through the meniscus, during normal gait can reach four times body weight and present significant challenges to fixation [20]. However, during unloaded knee motion the meniscus experiences only compressive forces [21,

22]. Therefore, fixation should maintain tissue approximation and neutralize shear stresses. For suture-based repairs, vertically oriented non-absorbable sutures are considered the gold-standard, because of load to failure [23]. This configuration encircles the strong circumferential fibers, maximizing strength. Meniscus repairs are weak at the scar after 12 weeks 24, and visual evidence of healing at second-look arthroscopy has been seen at up to four months [25].

Surgical Technique Of Meniscus

Fixation:

Arthroscopic techniques are the preferred method for meniscal repair; however no consensus exists as to the best technique. The most common indication for all-inside repair is tears of the posterior horn, as risk to neurovascular structures is decreased. All-inside repairs require less surgical time than other methods [26]. However, all-inside repairs do require an intact meniscal rim, highly specialized instruments, and implants. All-inside meniscal repair devices have progressed from rigid implants to current adjustable suture-based devices. Earlier versions of all-inside devices are no longer widely used or recommended. The adjustable suture-based all-inside devices are the state of the art.

Self-Adjusting Suture Containing Implants

The current generation of all-inside devices use ultra-high molecular weight polyethylene (UHMWPE) containing suture to connect typically non-absorbable poly ether-ether-ketone (PEEK) anchors.

The suture is pre-tied, typically with a sliding and self-locking knot. Insertion instruments require only standard anterior portals, and often use a disposable cannula or a skid to aid passage. The meniscus repair device is inserted through the inner meniscus fragment to a pre-determined depth of the peripheral rim, often guided by the cannula. Once both anchors are deployed, the sliding-locking knot is cinched to compress the tear. This adjustability allows appropriate tensioning for reduction and healing, and the option to place horizontal, oblique, or vertical configurations.

The FasT-Fix (Smith & Nephew, Andover, MA) was the first adjustable suture-based device (Fig. 1). It consisted of two 5 mm anchors, made of either poly L lactic acid (PLLA, absorbable) or polyacetal (nonabsorbable) connected by No. 0 non-absorbable braided polyester suture. The anchors are delivered by an instrument that is either straight or angled 22°. Once both anchors span the tear, the pre-tied sliding-locking knot is tensioned using a knot pusher/suture cutter. The original design was modified to become the Ultra Fast-Fix by reconfiguring the needle to facilitate insertion, and replacing the suture with a stronger No. 0 UHMWPE UltraBraid. The current iteration is the FasT-Fix 360 (Fig. 2), in which the anchors have been reconfigured to PEEK with an arrow design, and the suture is now No. 2-0 UltraBraid.

The RapidLoc (Mitek, Raynham, MA) was an adjustable suture-based device, consisting of a PLLA "backstop" and a

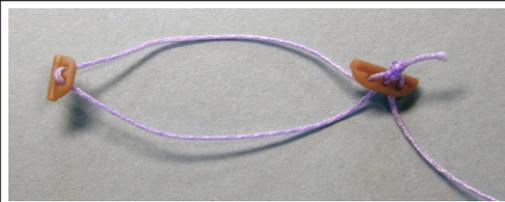


Figure 4: The OmniSpan has a strand of No. 2-0 OrthoCord doubled between two PEEK anchors. The sliding-locking knot is outside the first anchor, creating a repair with two sutures between the anchors with no knot on the meniscal surface. (copyright by author)

PLLA or polydioxanone (PDS) “top hat”, connected by either a No. 2-0 absorbable Panacryl or non-absorbable braided polyester suture (Fig. 3). The “backstop” anchor was placed across the tear to be extra-capsular, and the pre-tied sliding knot and “top hat” was then advanced, compressing the tear. The instrument included straight, 12° and 27° angled needles.

The OmniSpan (Mitek, Raynham, MA) replaced the RapidLoc, and uses a loop of No. 0 OrthoCord (55% PDS and 45% UHMWPE) suture between two PEEK anchors (Fig. 4). The sliding-locking knot is outside the loop, reinforcing the first anchor, and forming a double suture repair without a knot on the articular surface. Both loops of the repair are tightened concurrently, allowing equal tension. This device allows sutures to be placed in both horizontal and vertical mattress fashion.

The Meniscal Cinch (Arthrex, Naples, FL) has undergone incremental improvements since its inception (Fig. 5). The device is inserted with a 15° curved “gun” containing two separate trocar needles. It has an adjustable depth limiter on the handle, which is most commonly used at 18 mm. Each needle is loaded with a tubular PEEK anchor, and connected with a No. 2-0 FiberWire composed of UHMWPE and braided polyester (Arthrex Inc, Naples FL). The system includes a blue plastic “shoehorn” cannula to facilitate insertion, which is 6 mm in diameter and requires a large portal. The instrument allows placement of a vertical mattress stitch, secured with a pre-tied sliding-locking knot. After insertion, the first needle is removed and handed off. The second needle is “clicked” into position, and then a second

device is inserted. Once both devices are deployed, the suture is gently pulled at the handle to tension the repair. A disposable knot pusher/suture cutter is provided.

The Sequent meniscal repair device (ConMed Linvatec, Largo, FL) utilizes No. 0 Hi-Fi (braided UHMWPE) suture with up to seven PEEK anchors measuring 1.3 mm in diameter and 5.1 mm long (Fig. 6). Each anchor is placed individually through the meniscus, with a straight or 15° curved instrument, and deployed on the extra-capsular surface. The suture is then tensioned to set the anchor into the tissue, and additional anchors can then be placed with the same device. A minimum of 3 anchors must be inserted to complete the repair, although more can be used to create an all-inside continuous stitch. This allows numerous stitch configurations, from continuous to interrupted stitches, and vertical or horizontal mattresses. This is the only device that can place multiple stitches without removal from within the joint. However, the technique is demanding, and practice in the laboratory prior to use is advised. The set includes a side-loading disposable suture cutter for use at completion.

The MaxFire MarXmen (Biomet Sports Medicine, Warsaw, Indiana) is a self-adjusting all-inside all-suture implant with No. 0 MaxBraid PE (UHMWPE) and two braided polyester sleeves serving as anchors (Fig. 7). It is similar to the Juggernaut all-suture anchor in design and function, but modified for the meniscus. The instrument uses a needle (straight or curved) to insert the suture and two polyester anchors through the meniscus. The sliding-locking knot allows tensioning, and devices can be placed in either a horizontal or vertical



Figure 5:

- a) The Meniscal Cinch loaded in the insertion instrument, and
b) the device consisting of two PEEK anchors connected by No. 0 FiberWire, which uses a pre-tied sliding-locking knot. (copyright by author)

mattress fashion.

The CrossFix meniscal repair system (Cayenne Medical, Scottsdale, AZ) passes a No. 0 Force Fiber (UHMWPE) suture through two parallel 15 gauge hollow needles (straight or curved 12°, Fig. 8). Once the needles penetrate the meniscus, crossing the tear, a small shuttle passes the suture from one needle to the other on the extra-capsular surface. As the needles are withdrawn, a 3 mm horizontal mattress suture is left, and a pre-tied sliding Weston knot is advanced to secure the reduction. Additional arthroscopic knots can be added as reinforcement, if desired.

The AS (all suture) Repair device (Covidien, Minneapolis, MN) is similar to the CrossFix in design and function (Fig. 9). While the two needles are the same size, the AS repair device has conical solid needles with a polymer coat (NuCoat) to facilitate penetration. The instrument can be straight or curved 15°, and passes a No. 2-0 UHMWPE suture using a similar shuttle needle, but uses a modified Tennessee slider knot with two half hitches to secure the repair. Both instruments result in a 3 mm wide horizontal mattress, with a knot on the meniscal surface that risks chondral injury. Due to the instrument dimensions, only horizontal mattress sutures are possible.

Conceptually, this newest generation of all-inside suture based devices allows improved reduction, tissue compression, and stability compared to previous iterations. The overall goal of all-inside meniscal repair devices is to decrease complications seen with the earlier generations, and promote healing. However, these devices can generate significant tension which may be detrimental, leading to implant failure. Few



Figure 6: The Sequent has up to 7 PEEK anchors connected with No. 0 Hi-Fi suture, and allows continuous stitching technique. The stitch is secured at each anchor by two full clockwise rotations of the instrument, followed by tensioning to lock the suture into the anchor slot. (copyright by author)

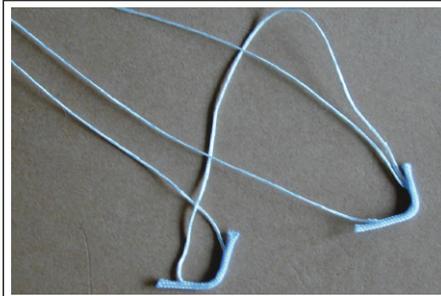


Figure 7: The MaxFire device is made of MaxBraid suture with two polyethylene sleeve anchors which bunch to create stability for fixation of the tear. (copyright by author)

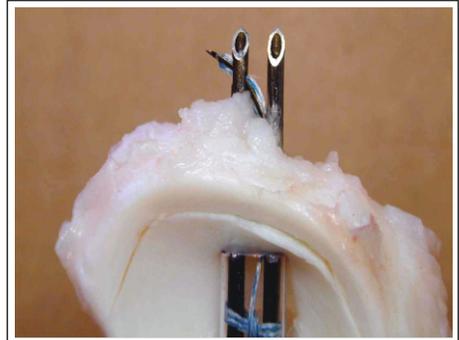


Figure 8: The CrossFix repair system passes a No. 0 Force Fiber suture through two parallel 15 gauge hollow needles. (copyright by author)

investigations regarding outcomes and complications of these adjustable all-inside suture implants are available in the literature.

Results Of All-inside Meniscal Repair:

Arthroscopic meniscal repair methods have similar outcomes to open methods, with the gold standard inside-out suture repair having a success rate of 82% [17, 27], and outside-in suture repairs having success rates as high as 87% [28]. The original suture-based all-inside technique described by Morgan reported good results, but without any long term follow-up [29]. Early devices which rely on arthroscopic knot tying demonstrate up to 90% success initially, but this declines to 81% at 1 year [30].

Suture based implants have good strength, and are biomechanically equivalent to the gold standard vertical mattress sutures [31,32]. However, as this is the latest generation of all-inside devices, there is little long term outcome data available. The RapidLoc has demonstrated success rates of 86 to 91% [33, 34, 35], but there are reported failure rates of 35% [36] with complications reminiscent of rigid devices [37]. Longer term follow-up of these devices shows re-operation rates of 48% [38]. The FasT-Fix has also shown success rates from 82 to 92% [39,40], but with limited reports of complications. All-Inside devices have been demonstrated to have greater failure strength than inside out alternatives in the repair of radial meniscal tears [41].

Comparison Of Meniscal Repair Devices:

Author's preferred technique:

A human cadaver knee based comparison of several all-inside meniscus repair devices was carried out by the senior author to compare the technical ease, reproducibility, and consistency of using these devices in human meniscus tissue. A needle penetration depth limited to 18mm was found to be anatomically safe. Curved needles effectively reached the posterior horn with minimal articular cartilage injury. However, significant differences were observed in the technical ease, reproducibility, and consistency of all these devices. The FastFix 360 and OmniSpan were easiest to insert, least likely to excoriate articular cartilage, and most consistent in performance. Yet, the OmniSpan did not have any knot or device on the surface to later damage the articular cartilage. Based upon this data the author's preferred technique uses the OmniSpan. The control provided by the gun allows for better positioning of the implants and decreased articular cartilage damage. Prototypes of the next generation of OmniSpan (the TrueSpan) perform even better but await clinical experience to confirm our expectations of superior performance.

Complications

All-inside meniscus repair has all the known risks and potential complications of knee arthroscopy. These occur in approximately 1% of patients, and include neurovascular injury, infection, and thrombophlebitis [42]. While neurovascular injuries are likely the most common complication of knee arthroscopy, when compared to other meniscal repair techniques the risk of this complication with all-inside repairs is

decreased. Neurologic injury rates as low as 2% have been reported for all-inside techniques, in comparison to 9% for inside-out repairs [43]. The development of the all-inside technique was primarily to eliminate the need for accessory incisions and suture passing that are responsible for most of the neurovascular risk, so that repairs in the posterior horn can be done more safely. Injury to the saphenous nerve is most frequent, but as it is a sensory nerve this is often of little consequence [44]. Peroneal nerve palsy and popliteal artery pseudoaneurysm have also been reported [45], as have cases of cyst formation and synovitis [46, 47].

Complications associated with the adjustable-suture based current generation of all-inside devices include over penetration of the implant, loss of fixation, inadequate tension, and problems with implant deployment [48,49]. An overall complication rate for all-inside repair of 19% has been reported comparable to the gold standard [43]. The RapidLoc has caused cartilage injury in limited reports [50,37], and cadaveric studies have demonstrated placement of these implants may be challenging, but the significance of this is unclear [48,49]. Complications of all-inside repair can be minimized with detailed knowledge of anatomy, proper portal placement, measurement of meniscal depth, and placement of the indicated implant in an appropriate and secure manner.

Rehabilitation:

Post-operative rehabilitation following meniscus repair is highly variable between surgeons, with little consensus in the



Figure 9: Covidien AS device uses two needles which are conical and solid needles, with a polymer coat (NuCoat) to facilitate meniscus penetration. (copyright by author)

literature. Early knee motion is thought to be advantageous, as prolonged immobilization is known to lead to stiffness, atrophy, and impaired healing of the meniscus [51]. However, higher degrees of knee flexion cause considerable posterior translation of the femoral condyles, which increases forces within the meniscus and may stress repairs [52]. Weight-bearing can help reduce and stabilize longitudinal (bucket-handle) meniscus tears due to radially directed hoop-stresses [22], but loads with knee flexion cause increasing shear forces in the meniscus. These forces are increased almost four times with the combination of weight-bearing and flexion to 90 degrees [52].

Based on this information, weight-bearing in full extension poses little risk to repairs of longitudinal meniscal tears, and

may aid with reduction and healing. However, for radial or meniscal root tear repairs (which are challenging with limited success), weight bearing is not advisable since circumferential fibers are not intact and the tear will be distracted. Accelerated rehabilitation programs designed to return patients to sport earlier have been described [53, 54], permitting early full

weight bearing and unrestricted knee motion. The only limitations on return to sport in accelerated programs are the resolution of postoperative effusion, and return of full motion. Thus far, results of accelerated programs have shown return to

sport without re-injury or complications.

Meniscal repair in the setting of ACL reconstruction presents unique challenges. There is no evidence to support slowing ACL rehabilitation for an associated meniscal repair, and with the increased stability of new adjustable suture-based devices there is less reason to do so [55]. The author's current protocol for modern all-inside devices allows immediate range of motion from 0 to 90 degrees, immediate full weight bearing, early closed-chain strengthening, flexibility and endurance training. After 2 months, full flexion is allowed, and full return sport is permitted once the knee has no effusion, has regained full extension, and demonstrates flexion to greater than 135°.

Conclusions

Advances in arthroscopy and instrumentation technology have made all-inside meniscal repair popular and effective in appropriate meniscal tears. While no arthroscopic method has proven to have superior outcomes in the literature, all-inside methods are indicated for posterior horn meniscal tears to minimize the risk to neurovascular structures. The

adjustable suture-based designs have so far demonstrated improved versatility and outcomes comparable to other methods. The versatility of these implants also allows their use in meniscal repairs that are not repairable by other methods, promoting the preservation of meniscal tissue when possible.

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