FiberStitch™ Implant

The FiberStitch implant is an innovative all-inside meniscal repair system that replaces hard PEEK implants with soft suture sheaths. 2-0 coreless FiberWire® suture and a pretied sliding knot provide secure arthroscopic all-suture meniscus repair. The ergonomic handle is designed for single-handed implant delivery and active implant deployment technology minimizes needle exposure beyond the meniscus, eliminating the need to past-point the needle.

Features and Benefits

■ Low-Profile Suture Implants
  • The low-profile suture anchors replace traditional hard PEEK plastic implants. Low-profile 2-0 coreless FiberWire® suture prevents tissue cut-through and minimizes friction against articular cartilage.¹

■ One-Handed Deployment
  • The ergonomic handle and easy implant deployment wheel allow true one-handed implant delivery.

■ Active Implant Deployment
  • The implants are deployed from the tip of the needle, reducing needle exposure beyond the meniscus.

■ Adjustable Depth Stop
  • The depth stop can be set with a single hand. Convenient markings in 2 mm increments allow setting adjustments from a minimum of 10 mm to a maximum of 18 mm.
Measure the meniscus using a measurement probe (1) or the laser lines on the tip of the FiberStitch needle (1a). Set the depth stop at a length 2 mm longer than the meniscus to ensure that the implant is fully deployed behind the meniscus.

The depth stop is set to 18 mm. To change the depth stop setting, engage the selector by depressing the gray tab (2) and the advancing it forward to the desired depth setting (2a).
Place the delivery needle over the desired entry point for the first implant. Advance the needle through the meniscus by pushing the entire handpiece forward until the desired depth is reached.

Roll the deployment wheel backward to a hard stop and an audible click is heard (4). This will deploy the first implant. The deployment wheel will display a solid vertical indicator (4a).
Roll the wheel forward to a hard stop and audible click. This will confirm that the first implant has been deployed. The deployment wheel will display 2 vertical indicators (a).

Remove the device from the meniscus and select the location to deploy the second implant. Advance the needle through the meniscus by pushing the entire handpiece forward until the desired depth is reached.

Deploy the second implant by rolling the deployment wheel backward to a hard stop and an audible click is heard. The deployment wheel will display a solid vertical indicator (a). Before removing the device from the tissue, roll the wheel completely forward to a hard stop (b).

Once the inserter is removed from the joint, a suture loop and a single suture will be visible outside of the joint. Insert a probe or similar device through the loop and pull the loop of suture to reduce the tissue. The suture spanning the 2 implants will reduce (a).  
**Note:** Continued tension on the loop suture after reducing the tissue will shorten the single suture. Only apply tension to the loop until the suture spanning the implants is taught.
Load the tail of the suture into the loading wire and pull the black tab to load the knot pusher.

Tension the single suture to reduce the suture loop. The implants are secured when the loop is reduced (a).

Advance the knot pusher towards the meniscus keeping it parallel to the suture while maintaining tension on the suture. Depress the back plunger to cut the suture (a).

Final construct.
Ordering Information

<table>
<thead>
<tr>
<th>Product Description</th>
<th>Item Number</th>
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<tbody>
<tr>
<td>FiberStitch™ Implant</td>
<td>AR-4570</td>
</tr>
<tr>
<td>Knot Pusher/Suture Cutter</td>
<td>AR-5815</td>
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<tr>
<td>Portal Skid</td>
<td>AR-4505</td>
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<tr>
<td>Meniscal Dart Probe</td>
<td>AR-4008</td>
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<tr>
<td>2.75 mm Suture Cutter, straight shaft</td>
<td>AR-11790</td>
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Products may not be available in all markets because product availability is subject to the regulatory approvals and medical practices in individual markets. Please contact Arthrex if you have questions about the availability of products in your area.

Reference

This description of technique is provided as an educational tool and clinical aid to assist properly licensed medical professionals in the usage of specific Arthrex products. As part of this professional usage, the medical professional must use their professional judgment in making any final determinations in product usage and technique. In doing so, the medical professional should rely on their own training and experience and should conduct a thorough review of pertinent medical literature and the product’s directions for use. Postoperative management is patient-specific and dependent on the treating professional’s assessment. Individual results will vary and not all patients will experience the same postoperative activity level or outcomes.

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