Medial Patellofemoral Ligament (MPFL)

Surgical Technique
The medial patellofemoral complex, consisting of the medial patellofemoral ligament (MPFL) and the medial patellotibial ligament, is the main passive stabilizer of the patellofemoral joint. Since it has been shown that rupture of the MPFL is the main pathological consequence of patellar dislocation\(^2\) and biomechanical studies have demonstrated that the MPFL is the main passive restraint against patellofemoral instability (PFI) and lateral patellar displacement, reconstruction of the MPFL has become a widely accepted technique for restoration of patellofemoral stability. Therefore, numerous techniques for reconstruction of the medial patellofemoral complex have been described with promising clinical results\(^3\). However, since it is known that a non-anatomic reconstruction of the MPFL can lead to nonphysiologic patellofemoral loads and kinematics\(^1\), the goal of surgical intervention must be an anatomic reconstruction. There have been multiple studies to evaluate the femoral insertion of the MPFL. Based on these anatomic\(^1\), biomechanical\(^6\) and radiologic\(^5\) findings, it is now possible to avoid the complications of increased patellofemoral pressure that is associated with non-anatomic (too anterior/proximal\(^1\)) fixation of the graft.

The Anatomic Double Bundle MPFL Reconstruction technique, developed in collaboration with Dr. Philip Schoettle from Zurich, Switzerland, replicates the native shape of the MPFL and provides the best possible stability in both flexion and extension. The Double Bundle technique also effectively limits rotation throughout the ROM minimizing postoperative instability. The technique, if accomplished directly and anatomically, may also provide for more aggressive rehabilitation protocols and earlier return to activity\(^4\).

As mentioned above, an important determinant of a successful outcome of MPFL reconstruction is the proper position of the femoral fixation of the graft, and our technique incorporates the use of a femoral template to ensure proper placement of the graft in the femur. This position provides a static fixation point that equalizes the tension across the graft in flexion and extension, thus minimizing the stresses across the patellofemoral joint.

**PATHOMORPHOLOGY OF PFI OVERVIEW**

The pathomorphology of PFI is dependent on different static and passive factors, such as alignment of the lower limb, dysplasia of the trochlea, and functionality of the MPFL. The patella is primarily stabilized by the medial patellofemoral complex from full extension to approximately 20° of flexion, and has no bony guidance, thus forcing the MPFL complex to bear the load of restraint against the lateralizing vector of the quadriceps muscle.

At about 20° of flexion the patella should engage into the trochlear groove, where the lateral trochlear facet is providing the static stabilization against patellar lateralization. The trochlea provides stability up to 60-70° of flexion, where the patella begins engaging into the notch. In cases of trochlear dysplasia, the patella cannot be guided properly and dislocation of the patella can occur more easily.

Very seldom, there are cases where the patella does not engage the notch in greater than 70° of flexion, and instability occurs. This can occur in cases of a valgus deformity or internal rotation of the distal femur, where the trochlear groove and the notch are positioned medially and the patella cannot engage. Chronic patellar dislocation is often seen in these cases with the patella tracking on the lateral condyle during the entire range of motion. In these cases, a realignment procedure should be considered.

**INDICATION FOR MPFL RECONSTRUCTION**

Since most cases of patellar instability occur in extension or slight flexion, with a slight underlying trochlear dysplasia, the majority can be treated with a reconstruction of the MPFL. The MPFL is ruptured in almost all cases after an acute patellar dislocation and is additionally weakened in cases of congenital trochlear dysplasia, since the patella tracks improperly from early childhood. The additional stresses and tension on the medial soft tissue complex from this maltracking can lead to an underdeveloped or insufficient MPFL and subsequent instability. Also, in cases with a PFI in deeper flexion, reconstruction of the MPFL should be considered as a concomitant procedure to provide stability in extension.
A gracilis autograft is used, as the size and strength has been shown to be sufficient for MPFL reconstruction (approximately 4 mm in diameter). Minimum graft length is 18 cm, whipstitched 10 mm at both ends. The graft should be tapered at each end to facilitate insertion of the graft into the patella.

**PREPARATION OF THE PATELLA**

Palpate the medial patellar border and make a 2 cm skin incision from the superomedial corner, extending to the center of the medial edge of the patella. Dissect down and expose the medial edge of the patella. Under fluoroscopic guidance, at a point 3 mm distal to the proximomedial corner of the patella, drill a 2.4 mm drill tip Guide Pin in a transverse fashion across the patella to a minimum depth of 25 mm. Place a second 2.4 mm Guide Pin 15-20 mm distal and parallel to the first one. Overdrill the 2 guide pins with a 4.5 mm cannulated reamer to a depth of 25 mm. (Note: Remove both guide pins after overreaming is complete.)
The proper position of the femoral insertion of the MPFL is very important to maintain proper biomechanics of the patellofemoral joint throughout the entire range of motion. The use of the MPFL template can help establish the position of the guide pin. The insertion point is approximately 1 mm anterior to the posterior cortex extension line (a), 2.5 mm distal to the posterior articular border of the medial femoral condyle (b), and proximal to the level of the posterior point of Blumensaat’s line. The template is placed on the area of the medial epicondyle on the distal femur and, under fluoroscopic guidance, a 2.4 mm guide pin (AR-1297L) is drilled across the femur and out through the lateral epicondyle. As the diameter of the doubled gracilis graft is predictably between 4 mm and 5 mm, the femur is drilled with a 6 mm Low Profile Reamer (AR-1406LP).

Pass the tails of one graft end through the eyelet of the first 4.75 mm SwiveLock™ and push the graft/anchor into the proximal drill hole until the eyelet is fully seated. Maintain tension on the suture limbs and screw the SwiveLock Anchor into the patella. After removal of the driver, the SwiveLock suture should be removed or tied to the graft sutures to reinforce the fixation. Repeat this procedure with the second graft end.

**PREPARATION OF FEMORAL INSERTION**
Loop the graft through the passing suture and pass the suture from the patellar origin to the insertion point at the medial femoral epicondyle. It is very important to maintain equal tension on both graft bundles during this step to ensure proper biomechanics of the patellofemoral joint. Deliver the sutures out of the medial incision, and, with equal tension on both graft bundles, pull the graft down to the medial epicondyle.

Drill to the far cortex. Maintain the 2.4 mm guide pin in the femur, as it will be used to pass the graft into the femur. Identify the space between the vastus medialis and the capsule, and bluntly dissect towards the femoral insertion area with a scissor, leaving the capsule intact. A right angle clamp is inserted into the prepared layer down to the medial epicondyle and the tip of the clamp is turned towards the skin. Make a 1 cm longitudinal incision and, using the clamp, pass a looped #2 FiberWire® back to the patellar insertion area.
FEMORAL GRAFT PASSAGE AND FIXATION

Place a 1.1 mm Nitinol guide wire into the drill hole next to the femoral guide wire to facilitate insertion of the 6 mm x 23 mm Bio-Interference Screw. Loop the graft sutures into the eyelet of the K-wire and deliver the suture tails out of the lateral femur prior to the graft entering the femoral socket. Using a clamp as a pulley, pretension the graft and insert it into the socket with equal tension on both graft bundles. Manually fixate the lateral patellar facet flush with the lateral femoral condyle with the knee at 30˚ of flexion. The isometry of the MPFL may be provisionally evaluated at this time by maintaining adequate tension on the graft and cycling the knee through the ROM. Once the proper isometry of the construct has been confirmed, final fixation can be accomplished. While maintaining tension on the graft, insert the 6 mm x 23 mm screw into the femur. Note: Evaluate the tracking and laxity of the patella throughout the knee ROM. If any adjustments need to be made, back out the femoral screw, make any tension or positioning adjustments and reinsert the screw.

POSTOPERATIVE CONSIDERATIONS

The Arthrex anatomical double bundle MPFL Reconstruction technique offers immediate, stable tendon-to-bone fixation and allows active quadriceps exercises between 0˚ and 90˚ of flexion at postoperative day one. A postoperative brace locked at 0˚-90˚ of flexion should be worn for a period of 6 weeks. Weight-bearing is limited to partial weight-bearing crutch ambulation until wound healing is complete and at that point can be increased according to the pain tolerance of the patient. Full ROM is allowed after 6 weeks with light jogging or cycling. The patient can resume full activity at 12 weeks.
ORDERING INFORMATION

*MPFL Convenience Pack (AR-1360B-CP) includes:*

MPFL Template

Bio-SwiveLock, 4.75 mm, qty. 2

Bio-Interference Screw, 6 mm

2.4 mm Guide Pin w/Suture Eyelet

Guide Pin, drill tip, 2.4 mm, qty. 2

Low Profile Reamer, 6 mm

Cannulated Drill, 4.5 mm


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.