Closure of Patellar Tendon Defect in Anterior Cruciate Ligament Reconstruction With Bone—Patellar Tendon—Bone Autograft: Systematic Review of Randomized Controlled Trials


Purpose: This study aimed to systematically review the highest level of evidence on anterior cruciate ligament (ACL) reconstruction with bone—patellar tendon—bone (BPTB) autografts with patellar tendon defect closure versus no closure after surgery. Methods: We performed a systematic review of multiple medical databases using Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. Level I and Level II randomized controlled trials comparing patellar tendon defect closure to no closure during ACL reconstruction with BPTB autografts were included. Two independent reviewers analyzed all studies. Descriptive statistics were calculated. Study methodological quality was analyzed using the Modified Coleman Methodology Score (MCMS) and Jadad scale. Results: Four studies with a combined 221 patients (154 male patients and 67 female patients) with an average age of 26.6 ± 2.4 years (range, 17 to 54 years) were included. All studies randomized patients before surgery into ACLR with BPTB autografts either with patellar tendon defect closure or without closure. There were no differences in clinical outcomes (Lysholm score, Tegner scale, International Knee Documentation Committee [IKDC] classification, modified Larsen score, and Lauridsen rating) between groups. There were no significant differences in knee pain between groups. All studies reported imaging findings of the patellar tendon defect, with 2 studies showing no difference in appearance between groups, one study showing excessive scar formation with defect repair, and one study showing improved restoration of normal tendon appearance with defect repair. The overall quality of the studies was poor, with all studies scoring less than 46 (average, 40.5 ± 4.7) on the MCMS and scoring 1 on the Jadad scale. Conclusions: Based on this systematic review of 4 randomized trials, there are no statistically significant or clinically relevant differences in outcomes between patients who have the patellar tendon defect closed and those who have it left open after ACLR with BPTB autografts. The methodology of the included studies limits the interpretation of the data, as evidenced by low MCMS and Jadad scores. Level of Evidence: Level II, systematic review of Level I and Level II studies.
patellar fractures, patellar tendon rupture, patellar subluxation, diminished extensor mechanism function, and anterior knee pain.\textsuperscript{15-19}

The surgical technique for ACLR with BTPB a autograft has been previously described\textsuperscript{7,8} and involves harvesting the central third of the ipsilateral patellar tendon with bone plugs from the patella and tibia. After incision through the skin and subcutaneous tissue, the peritenon surrounding the patellar tendon is incised, allowing full exposure of the patellar tendon, followed by harvest of the central third of the tendon. At the conclusion of the procedure, closure techniques are variable and can include closing both the patellar tendon and peritenon defects, closing either the patellar tendon or peritenon defect alone, or leaving both defects unrepaired. Currently, there is no consensus regarding the standard of care for management of the patellar tendon defect after ACLR with a BPTB autograft.

The purpose of this study was to review the published literature on ACLR with a BPTB autograft with patellar tendon defect closure versus no closure after surgery. We hypothesized that there would be no significant difference in clinical outcomes between patients undergoing patellar tendon defect closure compared with those without closure.

Methods

We performed a systematic review of multiple medical databases using Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.\textsuperscript{20} Before conducting the search, a systematic review registration was completed on August 12, 2013 using the PROSPERO International prospective register of systematic reviews (registration number CRD42013005357).\textsuperscript{21} Two independent reviewers (R.M.F., R.M.) completed the search, which was performed on August 13, 2013. The following databases were used: Medline (PubMed), CINAHL (Cumulative Index to Nursing and Allied Health Literature), and Cochrane Central Register of Controlled Trials. The following terms were searched: “anterior cruciate ligament,” “patellar tendon defect,” and “randomized.” The electronic search citation algorithm used was: acl[All Fields] AND (“patellar ligament”[MeSH Terms] OR (“patellar”[All Fields] AND “ligament”[MeSH Terms]) OR “patellar ligament”[All Fields] OR (“patellar”[All Fields] AND “tendon”[All Fields]) OR “patellar tendon”[All Fields]) AND defect[All Fields] AND closure[All Fields].

Inclusion criteria included English-language, Level I and Level II randomized controlled trials comparing patellar tendon defect closure to no closure after surgery. Exclusion criteria included non-English language studies, basic science or imaging studies, novel technique studies, scientific meeting abstracts/proceedings, and systematic reviews/meta-analyses. Levels of Evidence I and II were deemed inclusive (per the Oxford Centre for Evidence-Based Medicine used by the American version of the Journal of Bone and Joint Surgery\textsuperscript{22} and Arthroscopy). All references within included

![Diagram](https://example.com/diagram.png)

**Fig 1.** Systematic review search algorithm using Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines within Medline database. After application of all exclusion criteria, 4 studies were identified for final analysis.
<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Journal</th>
<th>Level of Evidence</th>
<th>Number of Patients</th>
<th>Average Age, yr</th>
<th>Age Range, yr</th>
<th>Number of Male Patients</th>
<th>Number With PT Closed</th>
<th>Number With PT Open</th>
<th>Number of Patients Available for Follow-up Duration, mo</th>
<th>Number of Patients Available for Follow-up (%)</th>
<th>No. of Right Knees</th>
<th>Graft Used</th>
<th>Surgical Technique</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adriani et al.</td>
<td>1995</td>
<td>KSSTA</td>
<td>I</td>
<td>61</td>
<td>26</td>
<td>17-52</td>
<td>40 (66%)</td>
<td>36 (59%)</td>
<td>25 (41%)</td>
<td>6</td>
<td>61 (100%)</td>
<td>33</td>
<td>BPTB autograft</td>
<td>Arthroscopically assisted ACLR with inside-out technique Side-to-side repair of tendon defect in 25 Tendon left open; peritenon closed in 36</td>
</tr>
<tr>
<td>Kohn et al.</td>
<td>1994</td>
<td>KSSTA</td>
<td>I</td>
<td>50</td>
<td>29</td>
<td>19-37</td>
<td>31 (62%)</td>
<td>25 (50%)</td>
<td>25 (50%)</td>
<td>30</td>
<td>40 (80%)</td>
<td>NA</td>
<td>BPTB autograft</td>
<td>Two-incision arthroscopically assisted ACLR with notchplasty Similar tunnel and screw sizes within groups Closed (group I): patellar defect packed with reamed bone ± additional cancellous bone from tibial head; peritenon closed with running No. 2-0 Vicryl sutures Open (group II): patellar defect loosely covered with gel foam, peritenon left open</td>
</tr>
<tr>
<td>Cerullo et al.</td>
<td>1995</td>
<td>KSSTA</td>
<td>I</td>
<td>50</td>
<td>23.5</td>
<td>17-34</td>
<td>43 (86%)</td>
<td>25 (50%)</td>
<td>25 (50%)</td>
<td>6</td>
<td>40 (80%)</td>
<td>NA</td>
<td>BPTB autograft</td>
<td>Closed group: 3 full-thickness simple interrupted No. 0 Vicryl sutures, followed by peritenon closure in same way if possible Open group: peritenon closure in same way if possible</td>
</tr>
<tr>
<td>Brandsson et al.</td>
<td>1998</td>
<td>KSSTA</td>
<td>I</td>
<td>60</td>
<td>28</td>
<td>17-48</td>
<td>40 (67%)</td>
<td>29 (48%)</td>
<td>31 (52%)</td>
<td>24</td>
<td>50 (83%)</td>
<td>NA</td>
<td>BPTB autograft</td>
<td>Two-incision arthroscopically assisted outside-in technique; meniscal pathologic process addressed as needed; notchplasty in 100%; paratenon closed in all with interrupted sutures Closed group: patellar tendon defect closed, patellar bone grafted Open group: neither of above</td>
</tr>
</tbody>
</table>

studies were cross-referenced for potential inclusion if somehow omitted from the initial search. Figure 1 shows the search strategy used according to PRISMA guidelines to generate the final study list.

After the original search, studies were reviewed for relevance by using previously described inclusionary and exclusionary criteria. Two independent reviewers (R.M.F., R.M.) analyzed studies deemed appropriate for inclusion. For all included studies, data collected included demographic data; intraoperative data, including type of graft, surgical technique, and method of defect closure (when performed); postoperative data, including rehabilitation, physical examination findings, pain level and other subjective outcomes, imaging findings (when available), return to activity, return to sport, reoperation rate, and complications. Study methodological quality was analyzed using the Modified Coleman Methodology Score (MCMS) and Jadad scale. Given the different methodology used in each of the studies, quantitative statistical analysis of the studies as a whole was not possible, and descriptive analysis was performed. Continuous variable data were reported as mean ± standard deviation. Weighted means and standard deviations were calculated for all participant, surgical, and outcomes parameters. Categorical variable data were reported as frequencies with percentages.

Results

Seven studies were identified with the initial search. Three of these studies were excluded, including one cadaveric study, one study that was not randomized, and one study that discussed topics unrelated to closure of the patellar tendon defect during ACLR with a BTPB autograft. Thus, a total of 4 studies met the inclusion criteria and underwent further analysis (Fig 1). These studies are described in detail in Table 1.

Table 2. Summary of Study Outcomes: Imaging

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Adriani et al.</td>
<td>26.91 (111.11)</td>
<td>37.54 (90.15)</td>
<td>16.47 (32.23)</td>
<td>30.5 (16.66)</td>
<td>“Binocular” appearance of 2 cords separated by a hyperechogenic bridge; with echogenic core surrounded by hyperechogenic ring; at 1 yr, echogenicity returned to normal; patellar bone defect evident at 1 yr</td>
<td>“Binocular” appearance of 2 cords separated by a hyperechogenic bridge; with echogenic core surrounded by hyperechogenic ring; at 1 yr, the cords still distinct with new tissue filling central area but different from true tendon structure; patellar bone defect evident at 1 yr</td>
<td>Patella alta ×1; patellar lengthening ×1</td>
<td>Patella alta ×1; patellar lengthening ×1</td>
</tr>
<tr>
<td>Kohn et al.</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>Normal appearance of PT after 2 yr; earlier healing of patellar defect</td>
<td>No healing of PT after 2 yr</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Cerullo et al.</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>No significant shortening</td>
<td>No significant shortening</td>
</tr>
<tr>
<td>Brandsson et al.</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>Central scar or gap at middle third with varying degree of tendon healing; lateral and medial thirds with oval-shaped hypertrophy (no difference between groups) Closed: complete healing in 11, partial healing in 5, no healing in 7</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
</tbody>
</table>

NA, not available; PT, patellar tendon; XR, radiograph.
All studies randomized patients before surgery into ACLR with a BPTB autograft either with patellar tendon defect closure or without closure. There were a combined 221 patients (154 male patients and 67 female patients) in the 4 studies. The average age of the patients was 26.6 ± 2.4 years (range, 17 to 54 years). The average postoperative follow-up was 16.5 ± 12.4 months (range, 6 to 30 months), with an average 85.8% ± 9.6% follow-up rate. A total of 52% (n = 115) of knees underwent patellar tendon defect closure, whereas 48% (n = 106) of knees did not.

Surgical technique, including closure of the patellar tendon defect, varied by study, with the most common technique involving side-to-side closure with Vicryl suture (Ethicon, Somerville, NJ). Adriani et al. performed arthroscopically assisted ACLR using an inside-out technique, followed by either closure of the patellar tendon defect with patellar bone grafting (n = 29) or no-closure/no-grafting (n = 31); in both groups, all patients underwent closure of the peritenon with interrupted sutures. Cerullo et al. performed arthroscopically assisted ACLR in all patients followed by either patellar tendon closure with 3 full-thickness simple interrupted No. 0 Vicryl sutures (n = 25) or no closure of the defect (n = 25); all patients underwent peritenon closure with interrupted Vicryl sutures as well. Kohn et al. performed 2-incision arthroscopically assisted ACLR using an outside-in technique, followed by either closure of the patellar tendon defect with reamed bone and peritenon closure with a running No. 2-0 Vicryl suture (n = 25) or loose coverage of the patellar defect with gel foam and no peritenon closure (n = 25). Only 2 of the 4 studies reported on their rehabilitation protocol, as illustrated in Table 1.

<table>
<thead>
<tr>
<th>Patellar Height on Lateral XR at 1 Yr: Closed</th>
<th>Patellar Height on Lateral XR at 1 Yr: Open</th>
<th>Patellar Shortening on XR: Closed</th>
<th>Patellar Shortening on XR: Open</th>
<th>Inferior Pole Spurs on PA-Frik at 6 Mo: Closed</th>
<th>Inferior Pole Spurs on PA-Frik View at 6 Mo: Open</th>
<th>Inferior Pole Spurs on PA-Frik View at 1 Yr: Closed</th>
<th>Inferior Pole Spurs on PA-Frik View at 1 Yr: Open</th>
<th>Computed Tomographic Findings at 6 Mo: Closed</th>
<th>Computed Tomographic Findings at 6 Mo: Open</th>
</tr>
</thead>
<tbody>
<tr>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
</tbody>
</table>

<2 mm: 14; 3-10 mm: 2; constant: 9

<2 mm: 14; 3-10 mm: 2; constant: 10

6 0 9 0 NA NA NA NA NA

N = 20

14 thickened 2x; 6 thickened 1.5x; width 15% less than normal; scar tissue throughout medial and lateral thirds

N = 20

15 normal; 5 slightly thickened; normal width; scar tissue central third

NA NA NA NA NA NA NA NA NA NA

NA NA NA NA NA NA NA NA NA NA

NA NA NA NA NA NA NA NA NA NA

NA NA NA NA NA NA NA NA NA NA
A summary of the physical examination findings, clinical outcomes scores, and imaging data is provided in Tables 2 and 3. Physical examination findings were inconsistently reported. One study commented on postoperative range of motion at 2 weeks after surgery, with nearly identical values in the open-defect (3.2° to 98.5°) and closed-defect (3.6° to 101°) groups. Also at 2 weeks postoperatively, the authors noted that 40% (n = 10) of patients in the open-defect group had pain while performing an isometric quadriceps contraction, compared with 56% (n = 14) with pain in the closed-defect group. Three of the 4 studies (75%) commented on postoperative inferior pole patellar pain, with essentially no differences between the open and closed groups. Pain ranged from 13% to 60% in the closed-defect group compared with 15% to 80% in the open-defect group.

Clinical outcomes scores, including Lysholm scores, Tegner scores, International Knee Documentation Committee (IKDC) scores, and modified Larsen and Lauridsen ratings, were variably reported. When provided, there were no differences in these outcomes between the defect-open and defect-closed groups in any of the studies. There were no significant differences in knee pain between groups, although one study noted increased painful spur formation in 36% of patients, which the authors attributed to bone grafting of the patellar defect. All studies reported on some form of imaging follow-up, including radiographs of the knee or advanced imaging (ultrasonography or computed tomography, or both) findings (or both) of the patellar tendon defect. Fifty percent (2 studies) found no difference in patellar tendon appearance between groups, 25% (one study) found excessive scar formation with defect repair, and 25% (one study) showed improved restoration of normal tendon appearance with defect repair.

Only one of the 4 (25%) studies reported on reoperation rates, with 7% (2 of 29) of patients undergoing reoperation in the closed-defect group, both for meniscal injuries. In the open-defect group, 6% (2 of 31) of patients underwent reoperation for either meniscal injury (one patient) and a recurrent traumatic ACL tear (one patient).

By definition, all 4 studies were considered Level of Evidence I or II. The overall quality of the 4 studies per the MCMS was poor, with all studies scoring less than 46 (average 40.5 ± 4.7). Similarly, all studies achieved a score of one on the Jadad scale.

**Discussion**

The principal findings of this study are as follows: (1) the data is inconsistent regarding the effect of patellar tendon defect closure on postoperative pain and function after ACLR with a BTPB autograft and (2) despite including only randomized clinical trials, the

### Table 3. Summary of Study Outcomes: Clinical

<table>
<thead>
<tr>
<th>Author</th>
<th>Larsen and Lauridsen Rating Scale Modified (Mean Score) at 6 Mo: Closed</th>
<th>Larsen and Lauridsen Rating Scale Modified (Mean Score) at 6 Mo: Open</th>
<th>ROM (°) at 2 Wk: Closed</th>
<th>ROM (°) at 2 Wk: Open</th>
<th>IKDC Score at 2 Yr: Closed</th>
<th>IKDC Score at 2 Yr: Open</th>
<th>Lysholm Score at 2 Yr: Closed</th>
<th>Lysholm Score at 2 Yr: Open</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adriani et al.</td>
<td>9.88</td>
<td>9.36</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Kohn et al.</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>No significant differences at 2 yr (only graphs; no actual numbers given)</td>
<td>No significant differences at 2 yr (only graphs; no actual numbers given)</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Cerullo et al.</td>
<td>NA</td>
<td>NA</td>
<td>3.6-101</td>
<td>3.2-98.5</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>Brandsson et al.</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>9 A, 13 B, 1 C, 1 D; no significant difference between groups</td>
<td>13 A, 11 B, 2 C, 0 D; no significant difference between groups</td>
<td>74 preoperatively to 95</td>
<td>76 preoperatively to 95</td>
</tr>
</tbody>
</table>

ACL, anterior cruciate ligament; IKDC, International Knee Documentation Committee; NA, not available; PT, patellar tendon; ROM, range of motion.
methodology of the included studies limits the interpretation of the data, as evidenced by the low MCMS and Jadad scores.

In the younger athletic high-demand patient population, ACLR with a BPTB autograft is the most commonly used graft choice. Even at the most competitive level with professional athletes, ACLR with a BPTB autograft is a reproducible surgical technique that uses a graft with high tensile strength, optimal fixation with bone-to-bone healing, and high return-to-play rates, with low overall complication and failure rates. Potential complications are rare, but dysfunction of the extensor mechanism, including patellar fracture, tibial tubercle fracture, or failure of the patellar tendon itself, remain the most worrisome. Lee et al. described a series of 1,725 consecutive ACLRs with BPTB autografts over a 20-year period and noted 3 complications related to patellar tendon harvest (0.2% complication rate), including 2 patellar fractures (one intraoperative and one postoperative) and one patellar tendon rupture (postoperative).

The critical contribution of the patellar tendon to the knee extensor mechanism, combined with the large central-third defect within the patellar tendon after a patellar tendon autograft harvest, certainly calls into question the role of patellar tendon defect closure after patellar tendon autograft harvest. Although ideally the decision to perform patellar tendon defect closure should be evidence based, given the paucity of data available in the literature, the intraoperative decision often comes down to balancing the potential perceived risks of closure (patella baja, decreasing patellar tendon length, suture irritation, increased operative time) with the potential perceived benefits (biological graft coverage by decreasing the large void created by an otherwise empty space left by harvest of the central third of the patellar tendon). Further, the relative individual contributions of bone grafting of the patellar defect versus closure of the patellar tendon defect versus closure of the peritenon on improving functional outcomes and decreasing postoperative pain are unclear.

Recently, Sobieraj et al. studied the mechanical implications of patellar tendon defect closure on the remaining patellar tendon after BPTB harvest. Using matched (by tendon dimension) pairs of fresh-frozen cadaveric patellar tendons, the authors harvested BPTB grafts from all the specimens and then performed defect closure in one half of the knees. After biomechanical testing, the authors noted no difference in load to failure, failure stress, stiffness, or modulus between the repaired tendons and those with the defect left open. In a separate cadaveric study, Eilerman et al. assessed the effect of patellar tendon harvest on patellofemoral contact pressures. The authors found no differences in patellofemoral joint pressures at varying

<table>
<thead>
<tr>
<th>Tegner Score at 2 Yr: Closed</th>
<th>Tegner Score at 2 Yr: Open</th>
<th>Local Tenderness to Palpation Above Inferior Pole: Patella Closed</th>
<th>Local Tenderness to Palpation Above Inferior Pole: Patella Open</th>
<th>Reoperations: Closed</th>
<th>Reoperations: Open</th>
<th>Main Conclusions</th>
</tr>
</thead>
<tbody>
<tr>
<td>NA</td>
<td>NA</td>
<td>60% at 6 mo</td>
<td>80% at 6 mo</td>
<td>NA</td>
<td>NA</td>
<td>No important differences between the groups Suturing peritendineum enhances healing and restores normal appearance of tendon; bone grafting patellar defect increases risk of painful spurs</td>
</tr>
<tr>
<td>NA</td>
<td>NA</td>
<td>32% at 5-8 mo</td>
<td>35% at 5-8 mo</td>
<td>NA</td>
<td>NA</td>
<td>No important differences between the groups, but closing tendon results in exuberant scar, so &quot;probably&quot; better to leave defect open</td>
</tr>
<tr>
<td>8 preoperatively to 6 postoperatively (significant worsening); no significant difference between groups</td>
<td>7 preoperatively to 5 postoperatively (significant worsening); no significant difference between groups</td>
<td>12.5% at 2 yr</td>
<td>15.4% at 2 yr</td>
<td>2 for meniscal injury within 2 yr postoperatively</td>
<td>1 for meniscal injury within 2 yr postoperatively; 1 for rerupture of ACL after new trauma</td>
<td>No important differences between groups; bone grafting patella and suturing tendon do not improve outcomes or reduce donor site morbidity</td>
</tr>
</tbody>
</table>

CLOSURE OF THE PATELLAR TENDON DEFECT IN ACLR 335
degrees of knee flexion (30°, 60°, and 90°) with and without side-to-side patellar tendon defect repair after central-third BPTB harvest. These cadaveric studies call into question the biomechanical role of patellar tendon defect closure.

In addition to biomechanical studies, several magnetic resonance imaging studies have found evidence of patellar tendon reconstitution after graft harvest. Nixon et al. reported that the size and intensity of the signal defect decreased over time, with an appearance identical to normal tendon tissue at 2 years after surgery. The authors performed biopsy procedures on 8 patients undergoing additional ipsilateral knee surgery after ACLR with BPTB autografts and noted essentially no histologic differences when compared with normal tendon histologic characteristics at 2 years after surgery. In a separate imaging study of 20 patients undergoing ACLR with BPTB harvest followed by defect closure, Coupens et al. found a nearly normal appearance of the patellar tendon at 1.5 years after surgery. Interestingly, the authors noted that throughout the follow-up period, the patellar tendon was found to have increased thickness compared with the contralateral leg but without any change in tendon width, despite the defect undergoing closure at the time of surgery.

Similar to the relatively inconclusive results found in these cadaveric and imaging studies, the results from the present systematic review are unable to provide evidence supporting or negating the routine use of patellar tendon defect closure after ACLR with a BPTB autograft. Although Adriani et al. noted increased scar formation on ultrasonography in the open-defect group compared with the closed-defect group at 1 year after surgery, their clinical, imaging and isokinetic findings were similar between the groups by 6 months after surgery. This finding led the authors to conclude that patellar tendon defect closure does not influence the extensor mechanism. Similarly, Brandsson et al. concluded that patellar tendon defect closure and bone grafting showed no improvement when compared with leaving the defect open based on finding no differences in pain, ultrasonographic findings, donor site morbidity, knee stability, or overall functional outcome at 2 years after surgery. In contrast, Cerullo et al. found computed tomographic evidence of a substantially thickened patellar tendon with scar tissue in the central third as well as the medial/lateral thirds in all patients undergoing defect closure compared with only 25% of patients in the open-defect group. Although there were no significant clinical or functional differences between the groups, the authors stated that it is "probably better" to leave the defect open after ACLR with BPTB autograft harvest. The results from the final study included in this systematic review further cloud the data because the authors found restoration of a normal tendon-like appearance with peritenon closure when compared with not closing the defect. Interestingly, the authors did find painful bone spur formation at the inferior pole of the patella in more than one third of patients who underwent patellar defect bone grafting, leading the authors to discontinue grafting while continuing with peritenon closure.

In a separate clinical study, Shaffer and Tibone used both intraoperative measurements and postoperative radiographs to determine the potential effect of patellar tendon defect closure on patellar tendon length and overall patellar position. In this study, 36 patients underwent ACLR with BPTB, with half of the patients undergoing patellar tendon defect closure and all patients undergoing peritenon closure. The authors found no evidence of clinically relevant patellar tendon shortening in the closure group and also found no evidence of patella baja. Other studies, however, have found evidence of patellar tendon shortening after ACLR with BPTB autografts.

Overall, the included studies represent the highest level of available evidence regarding the effect of patellar tendon defect closure on postoperative pain and function after ACLR with BPTB autografts. Nevertheless, it remains difficult to draw conclusions or make clinical recommendations based on this pooled data set. As noted, these studies did not assess kneeling pain or consistently measure any potential shortening of the extensor mechanism compared with the nonoperative knee, and this information would certainly be helpful in determining the clinical effects of patellar tendon closure. Similarly, it would have been clinically helpful for the patients in these studies to have subjectively assessed the presence or absence of pain along the anterior aspect of the knee postoperatively, especially with potentially provocative activities such as kneeling or stair climbing. As noted in a prospective study by Martin et al. in 1996, bone grafting the patellar defect significantly decreases patellofemoral pain compared with leaving the defect unfilled. In addition, a subjective assessment of the cosmetic appearance of the knee would have been interesting, because this type of data likely ties into overall patient satisfaction, an increasingly important entity in the current health care system.

The senior author (B.R.B.) has performed more than 2,200 primary and revision ACLR procedures with BPTB autografts and allografts. Our preference has been to graft the patellar and tibial tubercle defects with bone gathered at the time of tibial and femoral tunnel reaming. The patellar tendon defect is then loosely closed with the knee flexed at approximately 75° to 85° to reduce the likelihood of overconstraining the patella. The paratenon is subsequently closed with Vicryl suture. We have advocated patellar tendon defect closure
and bone defect grafting so that patients are not kneeling on the patellar defect or focused on the palpable defect in a nonclosed harvest site.

Limitations
The present study has several limitations. Because of the strict inclusionary criteria of including only randomized trials, the sample size is small with only 4 studies comprised of 221 patients. However, all studies included were of the highest possible level of evidence available on this topic. Despite using only Level I and Level II studies, the overall quality of the studies was low, as evidenced by the low MCMS and Jadad scores. Several of the studies omitted details from their methodology, which resulted in lower MCMS and Jadad scores; however, it is possible that those points were lost simply because of omission of details from the text of the manuscripts and not because of the quality of the actual studies. The major limitation is the lack of standardization between the studies with respect to the outcomes and imaging data collected, limiting the analysis to descriptive statistics and making comparisons between studies difficult.

Conclusions
Based on this systematic review of 4 randomized trials, there are no statistically significant or clinically relevant differences in outcomes between patients who have the patellar tendon defect closed and those who have it left open after ACL reconstruction with a BPTB autograft. Further, the data is inconsistent regarding the effect of patellar tendon defect closure on scar formation. The methodology of the included studies limits the interpretation of the data, as evidenced by low MCMS and Jadad scores.

References


44. Ferrari JD, Bach BR. Bone graft procurement for patellar defect grafting in anterior cruciate ligament reconstruction. *Arthroscopy* 1998;14:543-545.