Systematic Review of Arthroscopic Versus Open Repair for Recurrent Anterior Shoulder Dislocations

Jonathan Godin, MD, MBA* and Jon K. Sekiya, MD†§

Context: It remains unknown if arthroscopic repair of recurrent anterior shoulder instability is as effective as open repair.

Objective: The purpose of this study is to analyze the literature to provide clinical recommendations regarding the most appropriate therapeutic intervention for recurrent anterior shoulder instability.

Study Design: Systematic review of level I and II studies.

Data Sources: PubMed, EMBASE, the Cochrane Database of Systematic Reviews, and secondary references from 1967 to March 2010 were appraised for studies that met the inclusion criteria.

Study Selection: Inclusion criteria were English-language level I or level II trials involving the treatment of recurrent anterior shoulder instability. Exclusion criteria included non-English-language studies; level III, IV, or V studies; and trials examining treatment of first-time shoulder dislocation, posterior shoulder dislocation, or diagnoses other than recurrent anterior shoulder dislocations.

Data Extraction: Included studies underwent quality appraisal independently by each author identifying strengths, weaknesses, and biases.

Results: Four randomized controlled trials compared the use of arthroscopic and open repair for recurrent anterior shoulder dislocations. These studies show no statistically significant difference between the 2 operative approaches. No long-term follow-up data describing the effects of either surgical approach are available at this time. Each investigation had weaknesses in study design that decreased the validity of its findings.

Conclusions: While limited, the available evidence from randomized controlled trials does not show a statistically significant difference in redislocation rates, return to activity, and functional outcomes between the arthroscopic and open repair groups. Range of motion is marginally better following arthroscopic treatment when compared with open repair. Recommendations on the optimal surgical intervention cannot be provided.

Keywords: systematic review; recurrent anterior shoulder instability; open; arthroscopic; repair

The shoulder is the most commonly dislocated major joint, with a reported incidence of 1.7%. Symptomatic instability following dislocation is common, especially in young, active people. Recurrent instability, occurring in 50% to 90% of patients who first dislocate under the age of 20 years and in 40% to 74% of patients between the ages of 20 and 40 years, limits range of movement of the joint, requires multiple hospital and emergency department admissions for treatment, and often calls for surgical procedures to prevent further dislocation. Prior to arthroscopy, recurrent dislocations were managed by open repair, and the results of this approach, with only a 4% failure rate, were initially published by Dickson and Devas in 1957. There have been many studies documenting low recurrence rates ranging from 0% to 11% after open Bankart stabilization.

References 1, 3, 15, 17, 22, 38, 39, 41, 54-59.
The development of arthroscopic stabilization for recurrent anterior instability has undergone significant changes in the past 20 years. Possible benefits of arthroscopic stabilization include decreased length of hospital or outpatient surgery center stay, decreased postoperative pain, and improved range of motion (ROM). Initial arthroscopic fixation was performed by staple capsulorrhaphy, which resulted in recurrent instability in 16% to 33% of patients. Additional methods of arthroscopic stabilization have included transglenoid suturing, with a failure rate ranging from 0% to 49%, and bioabsorbable tack fixation, with a failure rate ranging from 9% to 23%. Newer techniques for arthroscopic stabilization have been developed, including suture anchor fixation and capsular plication, with failure rates ranging from 8% to 11%.

There are several reasons this systematic review is reasonable. Despite the proponents for both methods, it is unclear whether arthroscopic techniques equal the success of open techniques for the treatment of recurrent instability. There are a number of published reports directly comparing arthroscopic and open stabilization repairs.

Several meta-analyses and reviews of this topic have been conducted. However, each of these reviews included studies classified as level III evidence or lower. In contrast, this systematic review examines only level I and II trials. The high incidence of recurrent dislocation has implications for the individual and for society because chronic instability of the joint may prevent the individual from gaining employment or working at his or her potential. Moreover, with the growth in the number of orthopaedic surgeons specializing in shoulder surgery and sports injuries, as well as the advancement in arthroscopic techniques and sports medicine devices, there has been heightened interest in minimally invasive shoulder surgery for recurrent anterior instability. Thus, the comparison of arthroscopic versus open surgery for recurrent anterior shoulder instability is an area necessitating a scrupluous review.

**METHODS**

**Search Strategy**

A systematic review of level I and II trials between 1967 and 2010 was performed. PubMed, the Cochrane Database of Systematic Reviews, EMBASE, and secondary references were appraised for studies that met the inclusion criteria. Reference lists of retrieved articles were screened for additional publications. Specifically, the bibliographies of studies assessed for inclusion in this review, as well as periodicals focusing on the shoulder, were analyzed.

**Study Selection**

Publications included met strict selection criteria. Randomized controlled trials (RCTs) directly comparing arthroscopic stabilization with open repair for recurrent anterior shoulder instability were given priority, though level II evidence was included in the review. Patients were limited to adults (age 18 years or older) with recurrent anterior shoulder instability, which was confirmed by physical examination with a positive apprehension sign or increased anterior translation. Studies using outcome measures for shoulder function or pain—including recurrence (redislocation, subluxation, laxity), return to activity, Rowe and Constant scores, and ROM—were included. Studies with the following characteristics were not included: studies with no direct comparison of arthroscopic and open repair; non-English-language studies; level III, IV, or V studies; and studies examining treatment of first-time shoulder dislocation, posterior shoulder dislocation, or diagnoses other than recurrent anterior shoulder dislocations.

These criteria were used to independently select the relevant articles for this review by a reading of all titles and abstracts retrieved by the search strategy. The abstracts found from the search were reviewed for evidence of a direct comparison between open and arthroscopic stabilization of recurrent anterior shoulder dislocations. The 2 independent reviewers arrived at a consensus for including 4 studies in this review. The complete articles were then obtained for the selected abstracts, and a manual cross-reference was conducted. The final selected articles were independently critically appraised to classify the study design as a measure of the level of evidence.

**Methodological Quality Assessment**

All publications were assessed according to a methodological quality list for the assessment of RCTs (Table 1). The quality assessment list was modified and adapted to better fit this study. The requirement of blinding patients or health care providers to the intervention was excluded because this kind of blinding is not possible in this type of RCT. Each assessment criterion was graded as positive/yes, negative/no, or unclear. A quality score was calculated for the selected studies by summing the positive answers. Items D or F, or both, were answered only if C or E, respectively, or both, were scored negatively. The maximum attainable score was 9. Studies were considered to be methodologically high quality when at least 7 items scored positively, while a score of 4 to 6 was considered medium quality and 0 to 3 was low quality. The quality appraisal was then discussed between the authors, and consensus was reached regarding the strengths, weaknesses, and value of the included studies.

**Data Extraction**

Data on the study population, description and standardization of interventions, outcome measures, and results were extracted from the selected studies. First-time outcomes include the following: recovery defined as return to preinjury level of activity (sports or work), reinjury or recurrence (including subsequent surgery), subjective instability, results from validated shoulder rating scales. Secondary outcomes include...
A systematic review was performed because meta-analysis was not possible because of the diversity in outcome measures among the included studies and the differing presentations of data (median scores, mean scores, relative risk ratios). The results were summarized by means of a qualitative analysis.

**RESULTS**

**Search Results**

The PubMed search resulted in 34 citations. Six more citations were found through the Cochrane Database. Citation tracking identified no other studies.

**Excluded Studies**

The title or abstract, or both, was used to exclude 12 articles based on the aforementioned study selection criteria, and 22 were retrieved for a more detailed evaluation. Next, 4 reviews were excluded because they are Cochrane Database reviews or meta-analyses of RCTs on the topic. An additional 11 studies were excluded because they are not RCTs. A study by Mohtadi et al was excluded because it is an interim report of an ongoing trial examining multidirectional instability of the shoulder, as opposed to unilateral anterior instability. Finally, 2 additional studies were excluded because the reports are limited to abstracts. Consequently, 4 RCTs met our inclusion criteria.

**Data Extraction and Analysis**

There were difficulties in extracting data from the included studies, and there is little consistency in the outcome measures used in them. There were 218 patients enrolled in the selected studies (Table 2). Each trial directly compared arthroscopic and open repair for recurrent anterior shoulder instability. Each RCT-enrolled patient had unilateral, isolated, posttraumatic, recurrent anterior shoulder instability as verified by physical examination. In addition, Bottoni et al included patients experiencing both posttraumatic and atraumatic recurrent anterior shoulder instability.

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Table 1. Results of the methodological quality assessment for all included randomized controlled trials. A study by Mohtadi et al was excluded because it is an interim report of an ongoing trial examining multidirectional instability of the shoulder, as opposed to unilateral anterior instability. Finally, 2 additional studies were excluded because the reports are limited to abstracts. Consequently, 4 RCTs met our inclusion criteria.

| Item | Bottoni (2006) | Fabbriciani (2004) | Sperber (2001) | Jorgensen (1999) |
|------|---------------|------------------|--------------|-----------------|
| A. Was the treatment allocation randomized and concealed? | + | + | + | − |
| B. Was the outcome assessor blinded to the intervention? | + | − | − | + |
| C. Were the groups similar at baseline? | + | + | + | + |
| D. If not, were adjustments made in the analysis for differences of prognostic indicators at baseline and/or for confounding variables? | NA | NA | NA | NA |
| E. Was a sufficient proportion (≥80%) of included patients available for the full length of follow-up? | + | + | + | + |
| F. If not, was selective loss to follow-up excluded? | NA | NA | NA | NA |
| G. Was an intention-to-treat analysis included? | − | − | − | − |
| H. Were the interventions clearly defined? | + | + | + | + |
| I. Were the inclusion and exclusion criteria for study entry clearly defined? | + | + | + | + |
| J. Were the outcome measures suitable to measure clinically relevant differences in treatment effects? | + | + | + | + |
| K. Was the follow-up duration adequate to measure clinical differences between treatment modalities (≥1 year)? | + | + | + | + |
| Quality score (%) | 8 (88) | 7 (77) | 7 (77) | 7 (77) |

**Tables**

<references>8, 21, 23, 25, 29, 32, 34, 36, 52, 53, 59.</references>
Diagnostic arthroscopy was done prior to surgical randomization in 3 of the 4 trials. All 4 trials used suture anchors to repair the capsular laxity through the open technique (Table 3). However, there was variation in the arthroscopic stabilization method. Arthroscopic stabilization was performed using bioabsorbable tacks in the Sperber trial, while Jorgensen et al employed transglenoid sutures. Bottoni et al and Fabbriciani et al used suture anchors, thereby providing more consistent treatment regimens across the 2 groups.

Postoperatively, both treatment groups in all 4 studies were immediately immobilized with subsequent rehabilitation. The time spent in a sling varied among the trials, ranging from 3 weeks to 6 weeks. This difference in sling immobilization timing potentially affects the validity of the study’s results. Range of motion exercises were introduced after immobilization, and muscle-strengthening exercises were initiated thereafter in a consecutive manner in each study. Different regimens on the return to full external rotation and preinjury sporting activities were employed by the 5 trials. Bottoni et al allowed a return to sports or full active military duty at 4 to 6 months, depending on individual progress, while Fabbriciani et al allowed a return to sports activity after 6 months. Conversely, Sperber et al “discouraged overhead motion and contact sports for six months,” while Jorgensen et al allowed a return to sports activity after only 3 months.

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**Table 2. Patient demographics.**

| Study       | Total, No. | Sex, Men/Women | Age, y, Mean (Range) | Follow-up, mos, Mean |
|-------------|------------|----------------|---------------------|---------------------|
| Bottoni     | 32         | 29             | 25.2 (20-40)        | 28.5                |
| Fabbriciani | 30         | 26             | 25.0 (18-51)        | 36.2                |
| Sperber     | 30         | 26             | 24.5 (19-33)        | —                   |
| Jorgensen   | 21         | 20             | 28.0 (20-41)        | 36.6                |

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**Table 3. Surgical techniques (No.).**

| Study       | Transglenoid Sutures | Tacks | Anchors |
|-------------|----------------------|-------|---------|
| Bottoni     | —                    | —     | 32      |
| Fabbriciani | —                    | —     | 30      |
| Sperber     | —                    | 30    | 0       |
| Jorgensen   | 21                   | 0     | 0       |

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**Summary of Included Trials**

**Bottoni.** This is a level I trial conducted with a military patient population and a mean follow-up time of 29 months. Inclusion criteria included patients at least 18 years of age, subjective recurrent anterior shoulder instability, 6 months of failed rehabilitation, at least a 2+ anterior load shift test, and positive apprehension and relocation signs. Exclusion criteria included multidirectional instability, prior shoulder surgery, and less than 12 months at military assignment. Patients were randomized by sealed envelope, and participants wore T-shirts during postoperative physical examinations to aid blinding. Three patients were lost to follow-up, including 1 patient who was killed in military combat. Study results showed no statistically significant difference in ROM, similar failure rates between the 2 groups, and comparable functional outcome scores. The authors concluded that patients should be treated arthroscopically given the similar outcomes.

**Fabbriciani.** This is a level I trial conducted with a civilian patient population and a mean follow-up time of 24 months for isolated Bankart lesion using metallic suture anchors. Inclusion criteria included anterior shoulder instability secondary to trauma, no symptoms or shoulder surgery prior to trauma, no more than 4 episodes of anterior shoulder...
instability, Hill-Sachs lesion equal to or less than 30% of the humeral head surface, and a lack of multidirectional instability. Exclusion criteria included gross elongation or absence of the anteroinferior glenohumeral ligament, labrum detachment extended to the inferior part of the glenoid, anterior labrum periostial sleeve avulsion lesion, rotator interval tear, elongation or tear of the middle glenohumeral ligament, superior labral anterior posterior lesion, glenoid bone defect, and rotator cuff tear. Patients were randomized by sealed envelope, and the authors did not state the loss of any patients to follow-up. Study results showed no statistically significant Constant score difference, no recurrent dislocations in either group, ROM significantly greater in the arthroscopic group, higher Rowe score in the arthroscopic group (not statistically significant), no difference in pain between the 2 groups, nonstatistically significant increase in stability in the open group. The authors concluded that arthroscopic repair with suture anchors is effective for isolated Bankart lesions.

Jorgensen. This is a level II trial conducted with a mean follow-up time of 36 months in civilian patients with recurrent anterior shoulder dislocation. Inclusion criteria included a history of posttraumatic, recurrent, unilateral, anterior shoulder dislocation; complaints of shoulder instability that did not respond to a shoulder training program; normal ROM; positive apprehension sign; increased anterior translation; or negative sulcus test result on physical examination. Exclusion criteria included multidirectional instability or previous shoulder surgery. Patients were randomized by their address within Copenhagen, and participants wore T-shirts during postoperative physical examinations to aid blinding. No patients were lost to follow-up. Study results showed 1 arthroscopic group and no open group redislocations, 1 arthroscopic group subluxation and 2 open group subluxations, higher Rowe score in the open group (nonstatistically significant), higher Constant score in the arthroscopic group (nonstatistically significant), or greater ROM in the arthroscopic group. Given the results, in conjunction with fewer cosmetic problems and shorter hospital stays, arthroscopic repair for the treatment of recurrent anterior shoulder dislocation was recommended.

Sperber. This is a level I trial conducted with a mean follow-up time of 24 months. Inclusion criteria included patients at least 18 years of age; unilateral, recurrent, posttraumatic anterior shoulder dislocations or subluxations; and arthroscopically verified Bankart lesion. Exclusion criteria included bony Bankart lesion greater than 5 mm, primary dislocation, generalized joint laxity, bilateral instability, multidirectional instability, and soft tissue injury that could lead to instability. Patients were randomized by sealed envelope, and the authors did not state the loss of any patients to follow-up. Study results showed 7 arthroscopic group and 3 open group redislocations (nonstatistically significant), higher Rowe and Constant scores in the arthroscopic group (nonstatistically significant), and greater loss of external rotation in the arthroscopic group. They concluded that the arthroscopic repair with absorbable tacks resulted in less pain and minimal loss of external rotation but also a tendency toward a higher number of redislocations than in the open repair group.

Methodological Quality and Risk of Bias

The quality assessment scores of the 4 trials range from 7 to 8. According to our cutoff values for quality, all 4 trials were classified as high (Table 1); the 4 trials appear well conducted with minimum follow-up periods of 2 years. However, several methods utilized in the trials have the potential to inflict bias on the results. The relatively small sample sizes compared with the prevalence of the condition (218 total patients in the 4 studies combined) may yield random bias as a result of limited power. While treatment group allocation was randomized in all trials, it was not concealed in the Jorgensen trial. Patients were randomly assigned to arthroscopic or open treatment based on their address within Copenhagen. Blinding of outcome assessors was reported in detail in the Bottoni and Jorgensen studies, but it was not explicitly stated in the Fabbriciani and Sperber trials.

Another potential source of bias is the method used for data analysis. Intention-to-treat analysis was briefly mentioned in the Bottoni study, but it was not used in the other 3 trials. Patient baseline characteristics, percentage of patients available for follow-up, clinical appropriateness of outcome measures, and length of follow-up were comparable in all 3 trials. All the patients in the Bottoni study are active military personnel who may place more burden on their shoulders compared with the study participants enrolled from the civilian public in the other trials. This is a potential source of bias because it is not a true representation of the general population because of the rigorous physical demands placed on the cadets and their strict compliance with rehabilitation.

Primary Outcome Measures

Reinjury/recurrent instability. There was one redislocation in the arthroscopic group of the Jorgensen trial, while 1 patient treated arthroscopically and 2 patients treated with open repair in the Jorgensen trial suffered from postoperative subluxations. There were no recurrent dislocations in either group of the Bottoni or Fabbriciani studies. Two patients from the open repair group of the Bottoni study suffered reinjury. One patient had recurrent subluxation, and the other sustained a shoulder injury from an assault. There was no report of subluxation or reinjury in the Fabbriciani study. Two traumatic redislocations, 1 atraumatic redislocation, and 4 subluxations/reinjuries occurred in the arthroscopically treated patients from the Sperber trial. In the open repair group of the Sperber trial, there were 3 reinjuries, including 1 traumatic redislocation. It is not clear if any of the remaining 4 recurrences in the open group and the 2 other recurrences in the arthroscopic group were subluxations.
As a result of reinjury, 1 patient from the arthroscopic group in the Jorgensen trial underwent reoperation at 120 months. Three patients from the Bottoni study, all from the open repair group, underwent reoperation. Two patients in the arthroscopic group of the Sperber trial underwent reoperation, 1 for recurrent instability and the other to remove a broken Suretac tack. Similarly, the only reoperation in the open group of the Sperber trial was to remove a loose anchor. Fabbriciani et al did not report any reoperations.

Return to activity. Jorgensen et al employed a subjective patient survey to classify return to activity into 4 distinct groups. In the arthroscopic group, 18 patients (85.7%) regained good (7 patients, 33%) or excellent (11 patients, 52.4%) return to preinjury activity levels, while 2 patients (9.5%) were unchanged, and 1 patient (4.8%) suffered a reduction in activity level compared with pretreatment levels. Twenty patients (100%) treated by open technique in the Jorgensen trial regained activity, while 8 (40%) did so with some restrictions. Bottoni et al defined return to pre-injury activity as full, active military duty without physical limitations and restrictions. Fifty-seven patients (93.4%) returned to preinjury activity levels, while 1 patient in the arthroscopic group reported persistent shoulder pain without instability and was discharged from the military.

Functional assessment measures and shoulder rating scales. All 4 trials used the Rowe score, and none of them found a statistically significant difference in mean score between the arthroscopic and open repair groups. Three of the trials measured the Constant score without a statistically significant difference in mean score between the arthroscopic and open repair groups. In addition, Bottoni et al measured the Single Assessment Numeric Evaluation score, University of California–Los Angeles score, Simple Shoulder Test score, and Western Ontario Shoulder Instability score (Table 4). There were no statistically significant differences between the 2 treatment groups for any of these measures.

Pain. Persistent pain was reported with regard to treatment failure in the Bottoni trial. One patient from the arthroscopic group had persistent shoulder pain without instability and was medically discharged from the military. Fabbriciani et al reported pain as a subset of the Rowe and Constant scores. The study described identical results for the arthroscopic and open repair groups for both the Rowe score (8.5 ± 2.42) and Constant score (13.5 ± 2.42), thereby raising suspicion for the validity of the results. Sperber et al noted lasting pain in 3 participants, all of whom were in the open repair group. Fabbriciani et al did not specifically report data on pain.

Secondary Outcome Measures

Range of motion. None of the 4 trials found clinically significant differences in postoperative ROM between the 2 groups. Jorgensen et al found that 5 patients in the open group and 1 patient in the arthroscopic group had up to a 25% reduction in external rotation. In addition, 2 patients in the open group had up to a 50% reduction in external rotation and up to 25% reduction in abduction (P = 0.04). Bottoni et al found miniscule differences in 3 measures of ROM. The patients in the arthroscopic group exhibited statistically significant increases in external rotation compared with that of patients in the open group. However, the clinical ramifications and significance of these results are difficult to interpret. For example, Fabbriciani et al found a statistically significant difference between the 2 groups in the ROM portion of the Constant score, while Sperber et al found only a 1-degree-greater external rotation ROM in the arthroscopic group.

Objective instability. Jorgensen et al measured objective instability with the most analytic rigor. They incorporated an instrumented test with the DonJoy Knee Laxity Tester (Smith & Nephew) to assess anteroposterior translation of the shoulder joint. They reported a very small, statistically nonsignificant difference between the 2 groups. Objective instability was not

Table 4. Functional score outcomes.

| Study       | Rowe       | Constant   | SANE | UCLA | SST | WOSI |
|-------------|------------|------------|------|------|-----|------|
| Bottoni     | 91.6       | 86.0       | 93.5 | 90.6 | 94.4 | 90.0 |
| Fabbriciani | 91         | 86.5       | 89.5 | 86.7 | 90.0 | 95.0 |
| Sperber     | 100 (90-100)| 95 (75-100)| 100 (82-100)| 98 (67-100)| 95 | 90.8 |
| Jorgensen   | 92.5 (45-100)| 95 (55-100)| 62 (41-74) | 59.5 (30-71) | 433.6 | 505.8 |

*In means, with ranges in parentheses. SANE, Single Assessment Numeric Evaluation; UCLA, University of California–Los Angeles; SST, Simple Shoulder Test; WOSI, Western Ontario Shoulder Instability.*
reported in the Bottini or Fabbriciani trial. Sperber et al reported a positive apprehension test result in 3 patients treated with open repair, while none of the patients in the arthroscopic group had a positive apprehension test result.

**Patient satisfaction (subjective instability).** None of the studies explicitly reported patient satisfaction. However, subjective instability was assessed as a component of the Rowe score in the Fabbriciani and Jorgensen trials. The difference between the 2 groups in each study was not statistically significant.12

**Complications.** Neither Bottini et al nor Fabbriciani et al reported any perioperative complications. Sperber et al reported injury to the long thoracic nerve in 1 patient treated by arthroscopic intervention.51 Furthermore, 1 patient in each group of the Sperber trial underwent reoperation for broken surgical implants.51 Jorgensen et al reported 1 superficial wound infection and 1 patient complaining of hyperesthesia in the arthroscopic group.30 In addition, 4 patients in the open group complained of hyperesthesia around the scar.10

**DISCUSSION**

The 4 included trials each directly compare arthroscopic versus open treatment for recurrent, unilateral, anterior shoulder instability. The results displayed no statistically significant difference between the 2 groups with regard to recurrent instability, return to activity, and reoperation rates. For other outcomes, including shoulder function, either there were no statistically significant differences between the 2 groups, or the differences were small.

We identified 3 previously published systematic reviews with meta-analyses comparing arthroscopic and open repair for recurrent anterior shoulder instability.9,37,42 All 3 reviews combined studies with a variety of study designs and outcome measures, which can lead to heterogeneity in results.12

**Does open or arthroscopic repair lead to lower recurrence rates?** The RCTs included in this review demonstrated no significant difference in recurrence rates between patients treated by open technique and those treated by arthroscopic intervention.16,30,31 Two studies, Bottini et al and Fabbriciani et al, compared arthroscopic and open repair using suture anchors, which is the current gold standard surgical technique. These studies did not show statistically significant differences in recurrence rates between the 2 treatment groups.16 Meta-analyses reviewing this clinical question have drawn a different conclusion. Mohtadi et al documented a 2.04 odds ratio (P = 0.0003) of recurrent instability in those treated arthroscopically compared with those treated by open repair. Similarly, Lenters et al noted a 2.27 relative risk (RR; 0.00001) of recurrent instability, a 2.74 RR of recurrent dislocation (P < 0.0001), and a RR of 2.23 (0.002) for reoperation in patients undergoing arthroscopic stabilization compared with those treated by open technique. In a systematic review of 6 studies, Freedman et al reported that arthroscopic Bankart repair using transglenoid sutures or bioabsorbable tacks resulted in a higher rate of recurrence compared with open techniques (P = 0.01).50 It is important to note that each of these studies contained studies that are not level I or II evidence. In addition, surgical techniques varied by study, many of which are not currently accepted treatment methods.

**Is ROM superior following open or arthroscopic repair?** Arthroscopic intervention leads to a small improvement in postoperative ROM. Bottini et al and Sperber et al found no statistically significant difference in ROM between patients in the 2 treatment groups. Meanwhile, Fabbriciani et al and Jorgensen et al noted a statistically significant improvement in ROM along several axes following arthroscopic surgery when compared with ROM following open repair. These differences were most notable in external rotation.

**Does arthroscopic or open intervention lead to a better return to activity or function?** Bottini et al reported that 93.4% of patients had a sufficient return to activity level.4 However, 1 patient in the arthroscopic group was discharged from the military, while 3 patients in the open group were lost to follow-up. Jorgensen et al found that 85.7% of patients in the arthroscopic group and 100% of patients in the open group returned to preinjury activity levels.30 However, 7 patients in the arthroscopic group and 8 patients in the open group did so with some restrictions.30 In addition, Cole et al found that 14% of patients in the arthroscopic group and 18% of patients in the open group experienced moderate return to activity limitations, while 11% in the arthroscopic treatment group and 5% in the open repair group suffered from severe limitations in return to activity.4 Mohtadi et al noted that open repair offers a better outcome than arthroscopic repairs with respect to return to activity.4 However, the inclusion of studies with a wide variety of study designs and a high risk of bias limits the reliability of these findings. Lenters et al also found that open repair is superior with respect to postoperative functioning. They noted a RR of 0.87 (P = 0.03) for return to sports and work in the arthroscopic group compared with the open repair group.37

**Limitations**

A major limitation of this review is that only 4 level I or II RCTs comparing arthroscopic and open surgical repair have been conducted for this common shoulder injury. Because of this shortcoming, there is a potential for systematic bias in the validity of the evidence. Therefore, these findings should be interpreted with caution. During the review process, all of the qualified studies may not have been identified. The 4 RCTs vary in their inclusion and exclusion criteria, surgical techniques, suture material, number of tacks/suture anchors, and postoperative rehabilitation programs. Some results, such as ROM and return to activity, are difficult to compare because of differences in reporting throughout the studies.
Clinical Recommendations

SORT: Strength of Recommendation Taxonomy
A: consistent, good-quality patient-oriented evidence
B: inconsistent or limited-quality patient-oriented evidence
C: consensus, disease-oriented evidence, usual practice, expert opinion, or case series

| Clinical Recommendation | SORT Evidence Rating |
|-------------------------|----------------------|
| The recurrence rates following open and arthroscopic repair for recurrent anterior shoulder dislocations are similar. | A |
| Range of motion is slightly improved after arthroscopic treatment compared with open repair of recurrent anterior shoulder stability, and this outcome should be factored into clinical decision making. | A |

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