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A Randomized Comparative Study of a Standard Anterior Capsular Release versus Inferior Extended Release for the Treatment of Shoulder Stiffness

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Methods: Between January 2010 and December 2015, 39 patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups. In group I, 19 patients underwent arthroscopic capsular release of the rotator interval and anterior capsule. In group II, 20 patients underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval. The American Shoulder and Elbow Surgeons score, Constant scoring system, Simple Shoulder Test, visual analogue scale for pain, and range of motion (ROM) were used for evaluation before surgery, at 3, 6, and 12 months after surgery and on the last follow-up. **Results:** Preoperative demographic data revealed no significant differences (p > 0.05). The average follow-up was 16.07 months. Both groups showed significantly increased ROM at the last follow-up compared with preoperative (p < 0.05). At the last follow-up, no statistical differences were found (p>0.05) between groups I and II in functional scores and ROM (forward flexion, p=0.91; side external rotation, p=0.17; abduction external rotation, p=0.72; internal rotation, p=0.61). But we found that group II gained more flexion compared to group I at 3 months and 6 months (p < 0.05) after the surgery.

Conclusions: Both techniques of capsular release are effective for stiffness shoulder. However, the extended inferior capsular release shows superiority in forward flexion over anterior capsular release alone during 6 months of follows-up (level of evidence: Level I, therapeutic randomized controlled trial).

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Key Words: Shoulder; Stiffness; Frozen shoulder; Arthroscopic capsular release

Introduction

Shoulder stiffness is a common problem that is traditionally considered to be self-limiting.^{1,2)} Conservative treatment has usually been effective, with satisfying results reported in 60% to 80% of cases.³⁾ Many patients with rotator cuff tears not only have pain, but also have decreased range of motion (ROM) of the shoulder, which inhibits their function in daily life. Many factors can lead to such shoulder stiffness in patients with rotator cuff tears. The exact prevalence of shoulder stiffness combined with rotator cuff tears is unknown, but it is thought that mild shoulder stiffness, defined as at least a 20° of restriction in total ROM, exists in more than 40% of patients with rotator cuff tears.⁴⁾

Arthroscopic release of the glenohumeral joint has recently been used with success in the treatment of recalcitrant frozenshoulder syndrome.⁵⁻⁷⁾ Release of the contracture of the Coraco-

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Background: To compare the clinical outcomes of arthroscopic capsular release in patients with and without inferior capsular release for shoulder stiffness.

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humeral ligament (CHL) and thickening capsule is believed to be essential to treatment of frozen shoulder.^{7,8)} However, the extent of release is still the subject of debate. Besides the CHL and rotator interval, some authors have recommended release of the subscapularis tendon,⁹⁾ inferior capsule,¹⁰⁾ posterior capsule,^{11,12)} or global capsule¹³⁾ to improve elevation and internal rotation (IR), as well as external rotation (ER). Pearsall et al.¹⁴⁾ reported that information regarding preoperative motion loss would precisely direct selective capsular release.

A number of authors have also recommended posterior capsular release to improve IR.^{11,12)} However, according to Snow et al.¹⁵⁾ additional posterior capsular release was not associated with any significant difference in outcome when compared with anterior release. Chen¹⁶⁾ derived a similar result with extended posterior capsular release, showing that there was no advantage in function or ROM. Kim et al.¹⁷⁾ performed a level I study to compare anterior and inferior capsular release with anterior, inferior and posterior capsular release. In a one-year follow-up, there were no significant differences in ROM and functional scores between groups. Ranalletta et al.¹⁸⁾ suggested that isolated anteroinferior capsular release provides reliable improvement in pain and ROM for primary adhesive capsulitis.

The goal of this study was to determine the extent of capsular release needed in arthroscopic surgery for shoulder stiffness. To accomplish this, we compared the clinical outcomes of capsular release between those with and without inferior capsular release for shoulder stiffness. The hypothesis of our study was that the addition of inferior capsular release would contribute to improved outcomes.

Methods

Between January 2010 and December 2015, 152 consecutive patients with rotator cuff tear with concomitant stiffness were enrolled. Inclusion criteria were as follows: patients whose main symptom was shoulder stiffness, patients with small- to medium-sized full-thickness rotator cuff tear, and pain and functional limitation of the shoulder for at least 3 months. All patients underwent operation after the first visit to an outpatient clinic and had no conservative treatment period. There are no universally accepted criteria for the diagnosis of stiffness, but we determined stiffness by passive range of shoulder motion to be forward flexion of less than 100° (maximal 150°; forward flexion is glenohumeral motion without scapulohumeral rhythm), ER of less than 45°, or IR of the back at a level lower than the first lumbar spine junction.^{17,19)} Plain radiographs of the shoulder (true anteroposterior, supraspinatus outlet, and axial view) and shoulder magnetic resonance imaging were checked in all patients. Patients with concomitant glenohumeral pathologies (Bankart lesion, superior labrum anterior to posterior lesion), symptomatic cervical spinal lesion and large or massive cuff tears (tear size larger than 3 cm or more than 2 concomitant tendon tears) were excluded, as were patients with a previous history of operation or fracture. Overall, 55 patients were finally excluded. The baseline characteristics of the enrolled patients are shown in Fig. 1.

Patients were randomized into two groups through computer-generated blocked-randomization numbers (http://www. randomizer.org). After arthroscopic confirmation of inclusion and exclusion criteria, the surgical procedure was determined by a random number taken from a sealed envelope at the time of

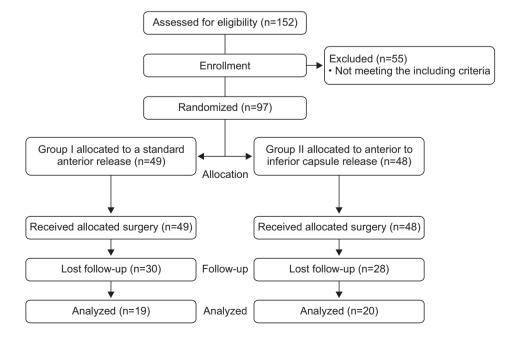


Fig. 1. Flow chart shows the conduct of the study according to Consolidated Standards of Reporting Trials (CONSORT) criteria.

surgery. All patients were blinded to this treatment and informed about the advantages and disadvantages of both treatments.

In group I, 19 patients underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release). In group II, 20 patients underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anteroinferior release). All patients gave informed consent before surgery and underwent the same rehabilitation protocol after the surgery. This study was approved by the institutional review board.



Fig. 2. Right shoulder glenohumeral joint viewing from the posterior portal. The rotator interval is done in the right shoulder glenohumeral joint until coracoacromial (CA) ligament and coracoid process base are exposed.

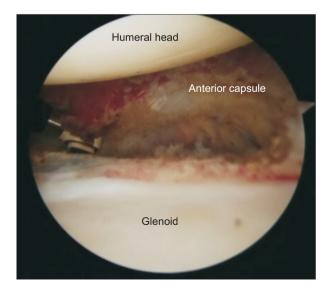


Fig. 3. Right shoulder glenohumeral joint viewing from posterior portal. Arthroscopic rotator interval release and anterior capsular release up to the 5:30 position in the right shoulder.



Fig. 4. Right shoulder glenohumeral joint viewing from trans-cuff portal. Arthroscopic rotator interval release, anterior capsular release and extended release of the inferior capsule up to 7:00 in the right shoulder.

Table 1. Preoperative Patient Demographic Data

Variable -	Group*		e volue
	Group I	Group II	<i>p</i> -value
Patient	19	20	
Age (yr)	62.32	60.05	0.41
Sex (men/women)	6/13	7/13	
Follow-up period (mo)	16.83	15.30	
Initial ROM (°)			
FF	130.00	121.5	0.61
ERa	78.42	76.00	0.35
ERs	73.16	68.50	0.30
IR	4.53	4.05	0.65
VAS for pain	4.75	5.35	0.41
ASES score	52.89	44.95	0.26
Constant scoring system	73.44	67.40	0.12
KSS	66.78	62.60	0.27
Initial tear size (cm)			
AP	1.26 ± 6.1	1.34 ± 5.7	0.65
ML	1.69 ± 5.2	1.66 ± 4.7	0.83

Values are presented as number only, mean only, or mean \pm standard deviation. ROM: range of motion, FF: forward flexion, ERa: external rotation (ER) at 90° of abduction, ERs: ER at side, IR: internal rotation, VAS: visual analogue scale, ASES: American Shoulder and Elbow Surgeons, KSS: Korean shoulder score, AP: anteroposterior, ML: mediolateral.

*Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release); group II, patients who underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anteroinferior release).

Surgical Technique

A senior shoulder specialist conducted all surgical procedures with the patient under general anesthesia. The patients underwent standard glenohumeral arthroscopy in the lateral decubitus position. In group I, capsular release began with the rotator interval via 3.0 mm 90° electrocautery (Arthrocare, Sunnyvale, CA, USA) through the anterior portal (Fig. 2). Anterior capsular release began below the biceps origin by preserving the glenoid labrum. The process began superiorly with resection of the supe-

Table 2. Comparison of Range of Motion

Maaauna	Gro	Group*	
Measure —	Group I	Group II	<i>p</i> -value
FF (°)			
Initial	130.00 ± 12.3	121.50 ± 11.4	0.61
3 months	129.12 ± 9.1	131.25 ± 10.1	0.96
6 months	138.24 ± 6.1	137.50 ± 5.2	0.85
12 months	142.73 ± 5.4	142.31 ± 7.5	1.00
Last F/U	143.75 ± 5.2	143.00 ± 6.3	0.67
ERa (°)			
Initial	78.42 ± 19.3	76.00 ± 18.2	0.35
3 months	78.24 ± 10.2	78.50 ± 12.7	0.80
6 months	80.00 ± 9.1	83.50 ± 6.1	0.22
12 months	86.36 ± 5.6	87.69 ± 7.2	0.61
Last F/U	85.83 ± 4.1	85.00 ± 4.2	0.77
ERs (°)			
Initial	73.16 ± 20.5	68.50 ± 14.8	0.30
3 months	74.12 ± 15.4	73.50 ± 13.2	0.82
6 months	75.29 ± 9.1	77.00 ± 11.7	0.73
12 months	80.91 ± 8.7	82.31 ± 6.4	0.61
Last F/U	84.17 ± 5.3	80.00 ± 7.1	0.20
IR (°)			
Initial	4.53 ± 2.4	4.05 ± 3.0	0.65
3 months	5.71 ± 1.7	7.10 ± 1.5	0.18
6 months	7.76 ± 2.5	8.00 ± 3.1	0.78
12 months	10.36 ± 2.6	9.92 ± 2.3	0.65
Last F/U	10.50 ± 1.6	10.00 ± 1.3	0.46

Values are presented as mean ± standard deviation.

FF: forward flexion, F/U: follow-up, ERa: external rotation (ER) at 90° of abduction, ERs: ER at side, IR: internal rotation.

*Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release); group II, patients who underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anteroinferior release). rior glenohumeral ligament and CHL. Release continued as far medial as the coracoid process until visualization of the verticaloriented fibers of the coracoacroimal ligament and conjoint tendon. Next, the middle glenohumeral ligament was released without damaging the subscapularis tendon, and then the anterior band of the inferior glenohumeral ligament was released up to the 5:30 position in the right shoulder (Fig. 3).

In group II, the same procedures were conducted, but there was extended release of the inferior capsule up to the 7:00 position until the posterior border of the inferior glenohumeral ligament (Fig. 4). The posterior capsule was not addressed. To avoid axillary nerve damage, capsular release was conducted just off the glenoid rim without violating the glenoid labrum. We believe that axillary nerve damage can be prevented if the

Table 3. Comparisons of Range of Motion Improvement by Period

Measure	Group*		. 1
	Group I	Group II	– <i>p</i> -value
FF (°)			
Initial to 3 months	-0.88	9.75	0.03
Initial to 6 months	8.24	16.00	0.03
Initial to 12 months	12.73	20.81	0.17
Initial to last F/U	13.75	21.50	0.06
ERa (°)			
Initial to 3 months	-0.96	2.00	0.43
Initial to 6 months	1.58	6.5	0.08
Initial to 12 months	7.94	11.69	0.41
Initial to last F/U	8.41	9.00	0.67
ERs (°)			
Initial to 3 months	0.96	5.00	0.28
Initial to 6 months	2.13	8.50	0.17
Initial to 12 months	7.75	13.81	0.23
Initial to last F/U	11.01	11.50	0.66
IR (°)			
Initial to 3 months	1.18	3.05	0.14
Initial to 6 months	3.24	3.95	0.50
Initial to 12 months	5.83	5.97	0.96
Initial to last F/U	5.97	5.95	0.98
V7-1	1		

Values are presented as mean only.

FF: forward flexion, F/U: follow-up, ERa: external rotation (ER) at 90° of abduction, ERs: ER at side, IR: internal rotation.

*Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release); group II, patients who underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anteroinferior release). electrocautery stays just off the glenoid rim. Moreover, electrical stimulation of electrocautery was helpful in detecting proximity to the axillary nerve before direct injury.

For small-sized (<1 cm) rotator cuff tears, a single-row repair was conducted, and a trans osseous equivalent repair (suture bridge technique) was performed for medium-sized (1–3 cm) tears. The selection of the surgical method was based on the location and shape of the torn tendon. Acromioplasty was performed for all type II and III acromions, along with the removal of sub-acromial spurs. For type I acromions, the acromial undersurface was smothered.

Rehabilitation

The same standardized rehabilitation protocols were applied to both groups postoperatively. Starting on the first day after the operation, pendulum circumduction was conducted, including gentle passive ROM exercise. An abduction brace was applied for 1 month after the operation. Pulley exercises were prescribed to increase the flexion after 1 month. When passive shoulder ROM was restored to 90%, isometric exercises in all planes were recommended. Thera-Band exercises, strengthening exercises for the muscles stabilizing the scapula, and advanced muscle strengthening exercises with dumbbells were taught 12 weeks after the operation. All listed procedures were recommended until the last visit after 12 months. No limit was imposed on use of the shoulder within a tolerable extent.

Outcome Measurement

Each patient was assessed before surgery and at 3, 6, and 12 months after surgery and at the last follow-up via the American Shoulder and Elbow Surgeons score²⁰⁾ and Constant scoring system. ROM including forward flexion, ER at side, ER at 90° of abduction, and IR of the treated shoulder was measured with a goniometer, and a visual analogue scale (VAS) for pain (0, no pain; 10, the most severe pain) was used to evaluate all patients at each visit. IR, measured in the sitting position, was evaluated by the tip of the thumb reaching the vertebral level. For analysis, the vertebral level was numbered serially: 0 for any level below the sacral region and 1 point added for each level above the sacrum. The postoperative cuff tendon integrity was assessed at

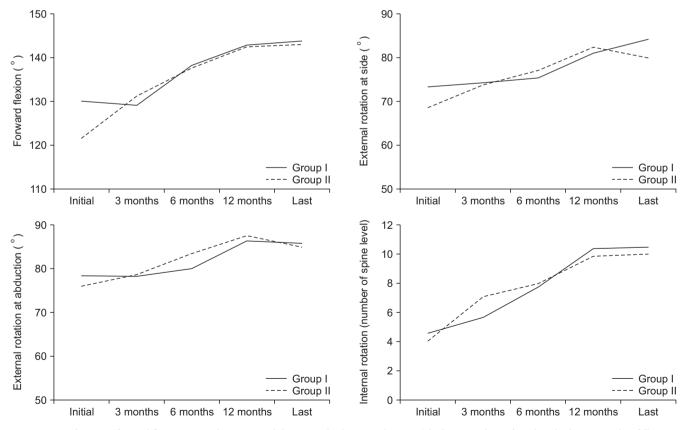


Fig. 5. Range of motion (forward flexion, external rotation at abduction and side, internal rotation) had improved significantly in both groups at last follow-up. There were no significant differences between the two groups at each time point. Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterrior release); group II, patients who underwent arthroscopic capsular release of the rotator interval (a standard anterinferior release).

8 weeks and 12 month after the operation using magnetic resonance imaging (MRI). All assessment data were prospectively collected by a clinical researcher who was blinded to the current study. Patients were also blinded during the assessment.

Statistical Methods

Statistical analysis was performed with IBM SPSS ver. 21.0 (IBM Co., Armonk, NY, USA). The Mann-Whitney test was used to compare differences between the outcomes of the 2 groups, and the Wilcoxon signed rank test was used to compare the differences in functional evaluation scores before and after surgery for each group. A p < 0.05 was considered significant.

Results

There were no significant differences in age or sex between groups (p > 0.05) (Table 1). Initial clinical scores before surgical release in both groups revealed no significant differences (p > 0.05) (Table 1). The mean follow-up period was 16.83 months for the standard anterior capsular release group (group

I) and 15.30 months for the extended inferior capsular release group (group II). In group I, there were 6 men and 13 women, with an average age of 62.32 years. In group II, there were 7 men and 13 women, with an average age of 60.05 years. On postoperative 12 months follow-up MRI, there was no significant difference in re-tear rate between groups. Three patients in group I (15.8%) and four patients in group II (20.0%) had type 4 and 5 re-tears according to the Sugaya classification (p>0.05).

The all ROM and functional scores improved significantly in both groups (p < 0.05) at final follow-up. In group I, the mean improvement of ROM was 13.75° in forward flexion, 11.01° in ER at side, 8.41° in ER at 90° abduction and 5.97 spinal levels in IR on the last follow-up. In group II, the mean improvement of ROM was 21.50° in forward flexion, 11.5° in ER at side, 9° in ER at 90° abduction and 5.95 spinal levels in IR at the last follow-up (Table 2).

We found that group II gained more flexion compared to group I from 0 to 3 months and from 0 to 6 months (p<0.05) (Table 3). However, no significant differences were found in other ROM, pain, and functional scores between groups at

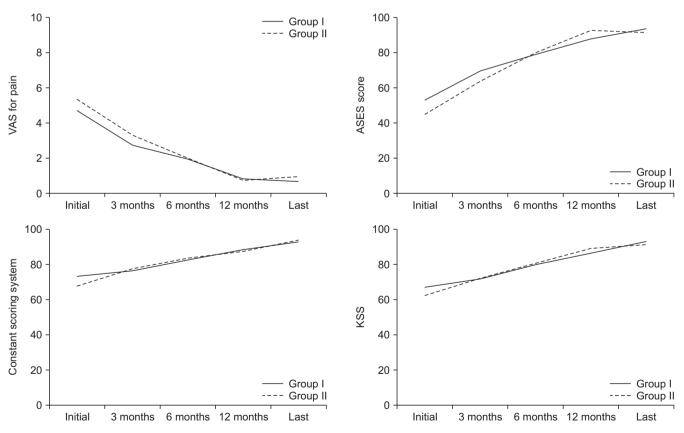


Fig. 6. Functional scores including visual analogue scale (VAS) score for pain, American Shoulder and Elbow Surgeons (ASES) score, Constant score, and Korean shoulder score (KSS) had improved significantly in both groups at last follow-up. There were no significant differences between the two groups at each time point. Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release); group II, patients who underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anterior release).

each time point (p>0.05) (Fig. 5, 6, Table 3, 4). No postoperative complications of infection, instability, or axillary nerve injury were observed in these patients.

Discussion

Shoulders with rotator cuff tears that were stiff before surgery were more likely to be stiff after surgery. A stiffer shoulder postoperatively correlated with better rotator cuff integrity post-

Table 4. Comparisons of Functional Scores

Variable	Group*		. 1
	Group I	Group II	<i>p</i> -value
VAS			
Initial	4.75	5.35	0.41
3 months	2.71	3.36	0.32
6 months	1.97	1.97	0.94
12 months	0.82	0.73	0.73
Last F/U	0.67	1.00	0.77
ASES			
Initial	52.89	44.95	0.26
3 months	69.29	63.39	0.28
6 months	79.12	79.50	0.94
12 months	88.09	92.55	0.16
Last F/U	93.33	91.70	0.76
Constant scoring system			
Initial	73.44	67.40	0.12
3 months	76.47	77.61	0.99
6 months	82.47	83.72	0.68
12 months	88.64	87.73	0.80
Last F/U	92.92	93.30	0.72
KSS			
Initial	66.78	62.60	0.27
3 months	72.18	71.78	0.64
6 months	79.38	80.67	0.71
12 months	86.45	88.82	0.55
Last F/U	92.92	91.50	0.63

Values are presented as mean only.

VAS: visual analogue scale, F/U: follow-up, ASES: American Shoulder and Elbow Surgeons, KSS: Korean shoulder score.

*Patients who underwent arthroscopic capsular release for shoulder stiffness were enrolled and randomized into two groups: group I, patients who underwent arthroscopic capsular release of the rotator interval, anterior (a standard anterior release); group II, patients who underwent arthroscopic capsular release of the anterior to inferior capsule, including the rotator interval (a standard anteroinferior release). operatively.²¹⁾ Therefore, concomitant rotator cuff repair with capsular release should be recommended to the patients who have rotator cuff repair with stiffness to achieve overall improved clinical outcomes.

Arthroscopic capsular release has become a reliable method for restoring ROM in patients with idiopathic frozen shoulder for which physical therapy and conservative care have failed.⁵⁾ A nearly normal ROM and good outcome scores can be obtained with this procedure. In the current study, ROM significantly improved compared with preoperative values in both groups; however, it showed superiority for group II in the forward flexion until 6 months after surgery.

Histologic studies, open exploration, and arthroscopic observations have revealed that the CHL and rotator interval are the major affected areas in frozen shoulder and should be released to restore passive ER.^{7,22-24)} Therefore, it was believed that the anterior capsular structures were required to be released to restore ER and abduction.^{14,25)} However, none of the studies conducted to date discussed the extent to which the anterior structures need to be released. In addition to the rotator interval, CHL, and middle glenohumeral ligament, Pearsall et al.⁹⁾ recommended releasing the intra-articular subscapularis tendon without significant morbidity.

A release of the superior and middle glenohumeral ligaments, the rotator interval, the CHL, and the intra-articular portion of the subscapularis increased the range of ER, whereas release of the anteroinferior capsule, including the anterior band of the inferior glenohumeral ligament, improved the elevation range. However, arthroscopic posterior capsular release is controversial. Selective posterior capsular release or circumferential release around the joint capsule was advocated in some studies for recovery of IR.^{6,20)} Kim et al.¹⁷⁾ found that the range of ER tended to be better in the posterior extended capsular release group until 6 months after surgery, but that it did not shown any significant differences and that observed differences did not last beyond one year after surgery. Moreover, Codding et al.²⁶ reported that selective arthroscopic posterior-inferior capsular release may be a reasonable solution in overhead athletes with symptomatic glenohumeral IR deficit.

We believe that treatment should be directed at the rotator interval and the contracted CHL, as this is the site of primary pathology in idiopathic adhesive capsulitis. We found that the range of elevation tended to be better in group II (standard anterior inferior capsular release) until 6 months after surgery. We cautiously state that there was an apparent benefit in terms of flexion with additional inferior capsular release. Miyazaki et al.²⁷⁾ found that there were better improvements in pain, IR and ER in patients who underwent anteroinferior capsulotomy.

It should be noted that there were several limitations to this study. First, the size of the enrolled patients is quite small. Moreover, there was no control group. For these reasons, it was difficult to determine what percentage of patients would have improved without rotator interval release and capsulectomy. Second, the mean follow-up period was relatively short. For both groups, it was 16.1 months. However, shoulder stiffness does not recur once it has healed, and it is unlikely that the results may get worse with time. Finally, there were no long-term results. Overall, the results might have been more reliable with more enrolled patients and additional long-term data.

Conclusion

Arthroscopic capsular release is a reliable method for relieving pain and improving clinical function in patients with shoulder stiffness. However, the addition of inferior capsular release shows superiority in forward flexion over anterior capsular release alone at 6 months follows-up.

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