Introduction

Irreparable posterosuperior rotator cuff ruptures in the young patient remain a challenging clinical entity. Owing to high revision rates reported in young patients ranging between 18 and 48%, reverse shoulder arthroplasty is considered the last-line treatment [6, 8, 17, 23]. Thus, the general clinical goal is to delay arthroplasty for as long as possible. Many therapeutic options including debridement, long head of biceps tenotomy, partial rotator cuff repair, tendon transfer [7, 14, 16], and subacromial spacer implantation [12, 28] have been proposed and performed with varying short- and often disappointing long-term results. Superior capsular reconstruction (SCR) was popularized by Mihata et al. [22] in 2012 to overcome superior humeral translational instability after massive rotator cuff tears. Originally, a fascia lata autograft harvested via a minimally invasive lateral approach from the ipsilateral side was used as a graft. Grafts with a thickness of 8 mm were initially used for the procedure [13]. In a biomechanical laboratory setting, it was shown that a significant reduction in superior humeral translation and an increase in subacromial distance could be achieved with reconstruction of the superior glenohumeral joint capsule, while these effects were enhanced by side-to-side suturing between the infraspinatus and the posterior margin of the graft [20, 21]. These promising in vitro results were then confirmed in a clinical study showing relatively quick pain reduction, followed by a significant increase in range of motion [19].

Owing to the morbidity associated with harvesting of fascia lata autograft, a human dermal allograft patch was introduced to overcome donor-site morbidity. Recent publications showed similar promising short-term results with significant reduction in pain and restoration of function after 1 year [2, 4, 11, 25]. However, whether this technique may be indicated as a treatment option for patients with pseudoparalysis remains unclear. Thus, the objective of this study was to investigate the functional and radiological short-term outcome after arthroscopic SCR with a human dermal allograft for partially irreparable rotator cuff tears. Furthermore, differences in outcome between patients with and without limited shoulder function were analyzed.

Patients and methods

All patients with irreparable supraspinatus tendon tear and intact or repairable subscapularis and repairable or partially repairable infraspinatus (ISP) tendon, with reduced daily activity and persistent night pain without improvement after conservative treatment including physiotherapy, who presented at our clinic in 2016 were eligible for arthroscopic SCR. All patients with a magnetic resonance image (MRI)-verified rotator cuff tear with retraction of Grade 3 according to Patte et al. [24] were considered for inclusion in the present study. Patients with an intraoperatively irreparable supraspinatus tendon tear were ultimately included in the study. Exclusion criteria were patients with reparable supraspinatus tendon tears, irreparable infraspinatus tendon and/or subscapularis tendon tears, and cuff tear arthropathy (>Grade 2 Hamada; [10]). Patients were divided into two groups depending on their preoperative range of motion. Patients with forward flexion over 90° were included into group 1. Patients with a forward flexion of 90° or below were considered as having pseudoparalysis and were assigned to group 2.

Of 45 initially enrolled patients, a reconstruction of the supraspinatus tear was possible in 24 patients, who were therefore excluded from the study, while 21 underwent arthroscopic SCR with a dermal allograft. The study was ap-
Table 1  Patient’ characteristics

|                      | Total (n = 21) | Forward flexion >90° (n = 9) | Forward flexion ≤90° (n = 12) | p      |
|----------------------|---------------|------------------------------|------------------------------|--------|
| Age (years)          | 65.9 ± 7.4 (50–77) | 67.2 ± 5.9 (57–75)            | 64.8 ± 8.4 (50–77)            | 0.475  |
| Gender (female/male) | 14 (67)/7 (33) | 7 (78)/2 (22)                | 7 (58)/5 (42)                | 0.642  |
| Shoulder side (right/ left) | 18 (86)/3 (14) | 9 (100)/0                   | 9 (75)/3 (25)                | 0.229  |
| Dominant arm involved (yes/no) | 20 (95)/1 (5) | 9 (100)/0                   | 11 (92)/1 (8)                | 0.999  |
| Smoker (non/ex/ current) | 14 (67)/2 (9)/5 (24) | 6 (67)/1 (11)/2 (22)         | 8 (67)/1 (8)/3 (25)          | 0.971  |
| Diabetes mellitus (yes/no) | 2 (10)/19 (90) | 1 (11)/8 (89)               | 1 (8)/11 (92)                | 0.999  |
| Prior surgery (yes/no) | 7 (33)/14 (67) | 1 (11)/8 (89)               | 6 (50)/6 (50)                | 0.159  |

RC tear characteristics

|                      |                      |                  |                  |       |
|----------------------|----------------------|------------------|------------------|-------|
| SSP tear             | 3 (14)               | 1 (12)           | 2 (17)           | 0.792 |
| SSP + SSC tear       | 1 (5)                | 0                | 1 (8)            |       |
| SSP + ISP tear       | 8 (38)               | 4 (44)           | 4 (33)           |       |
| SSP + SSC + ISP tear | 9 (43)               | 4 (44)           | 5 (42)           |       |
| RC repair with SCR   |                      |                  |                  |       |
| Patch repair         | 5 (24)               | 2 (22)           | 3 (25)           | 0.969 |
| Patch + SSC repair   | 3 (14)               | 1 (12)           | 2 (17)           |       |
| Patch + ISP repair   | 7 (33)               | 3 (33)           | 4 (33)           |       |
| Patch + SSC + ISP repair | 6 (29)             | 3 (33)           | 3 (25)           |       |
| Revision surgery indi cated | 6 (29)             | 2 (33)           | 4 (67)           | 0.659 |

ISP infraspinatus, RC rotator cuff, SCR superior capsular reconstruction, SSC subscapularis, SSCP supraspinatus

proved by the institutional review board (201609_EK13). Informed consent was obtained from all patients.

Surgical technique

All surgical procedures were performed by the same surgeon. The surgical procedure has been described in detail in previous publications [18, 25]. In short, the patient was operated on in the beach chair position starting with a standardized diagnostic arthroscopy. The long head of biceps tendon, if still present, was tenotomized at its origin. Subacromial soft tissue debridement and an acromioplasty were performed in all cases. If the subscapularis tendon showed a tear, an arthroscopic repair was performed. Repairable tears of the infraspinatus tendon were addressed with partial repairs in order to re-create the force couple between anterior and posterior rotator cuff. The superior capsule was then reconstructed using a decellularized human dermal allograft (ArthroFLEX®, Lifenet Health, Virginia Beach, VA, USA). The graft was fixed to the glenoid in standardized fashion using two suture anchors (Arthrex Corkscrew®—FT3, Arthrex, Naples, FL, USA). The arm was then brought into 30–45° of abduction and the graft was fixed to the humeral head using a knotless transosseous equivalent repair configuration with high-strength suture tapes (SpeedBridge™ and FiberTape®, Arthrex, Naples, FL, USA). Three side-to-side sutures between the infraspinatus and the dorsal part of the graft were performed in 5–7-mm distance. No closure of the anterior rotator cuff interval was performed in any of the cases.

Postoperative treatment and rehabilitation

Postoperative treatment was standardized for all patients. After surgery, patients were immobilized in a sling for 6 weeks. Only passive mobilization exercises were allowed. Assistive and active physiotherapy were allowed after 6 weeks with no strengthening exercises or load-bearing for 12 weeks postoperatively.

Radiological, clinical, and subjective assessment

Magnetic resonance imaging of the shoulder was performed before and 11.6 ± 0.6 months (range, 11–13 months) after arthroscopic SCR and images were evaluated by two orthopedic surgeons independently (M.E., P.H.). Evaluation of the MRI was performed in standardized fashion using frontal, parasagittal, and coronal axis slices on a 3-Tesla MRI unit. The Structural integrity of the SCR and repaired rotator cuff tendons was evaluated using Sugaya's MRI classification considering Grades 1, 2, and 3 as intact, and Grades 4 and 5 as re-torn [29]. Tendon retraction was graded according to Patte et al. [24] Clinical and subjective assessment was made at baseline as well as 4.2 ± 1.5 months (range, 3–7) and 11.6 ± 0.6 months (range, 11–13) postoperatively. Shoulder function was assessed using the total Constant score (CS) and its subgroups pain, activity of daily living (ADL), range of motion (ROM), and strength. Subjective evaluation included the American Shoulder and Elbow Surgeons score (ASES, 0–100 points) and the Simple Shoulder Test score (SST, 0–12 points). Additionally, outcome was compared between patients with (abduction and/or flexion of 90° or below, group 1) and without (abduction and/or flexion over 90°, group 2) pseudoparalytic conditions [30]. An MRI-verified rupture of the dermal graft was considered as a study endpoint (treatment failure). Complications occurring during the study period were recorded.
Abstract · Zusammenfassung

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M. Eigenschink · L. Pauzenberger · B. Laky · W. Anderl · R. C. Ostermann · P. R. Heuberer

Arthroscopic superior capsular reconstruction using a human dermal allograft in patients with and without preoperative pseudoparalysis

Abstract

Background. Massive irreparable rotator cuff tears (RCT) may cause severe functional impairment and pain as a result of loss of superior humeral stability. Reconstruction of the superior capsule (SCR) has been reported to restore glenohumeral stability and function.

Objective. The purpose of this study was to investigate short-term functional and radiological outcome after arthroscopic SCR with a human dermal allograft for irreparable RCT.

Methods. In total, 21 patients (mean age 65.9 years, 50–77), who underwent arthroscopic SCR were prospectively enrolled in the present study. Magnetic resonance images (MRI) were obtained before and 12 months after surgery to evaluate graft survival. Functional and subjective outcome including the Constant score (CS), the American Shoulder and Elbow Surgeons score (ASES), and the Simple Shoulder Test (SST) was evaluated preoperatively as well as 3–6 and 12 months postoperatively and was compared between patients with and without preoperative pseudoparalysis. Complications were recorded.

Results. The MRI evaluation revealed intact grafts in 71.4% of patients 1 year postoperatively; these patients showed significant improvements from baseline to follow-up time points regarding the total CS and its subgroups pain, activity of living, range of motion, strength, ASES, and SST (p < 0.01). Pseudoparalysis was present in 57.1% of cases preoperatively, but was reversed in 100% of cases with intact graft at the 1-year follow-up. The outcome was similar between groups at final follow-up. Complications occurred in 29% (one infection, five graft failures) of patients and were indications for treatment with reverse shoulder arthroplasty.

Conclusion. Arthroscopic SCR with a human dermal allograft in patients with irreparable RCT restored shoulder function and relieved pain in patients without and with preoperative pseudoparalysis.

Keywords

Pseudoparalysis · Superior capsular reconstruction · Massive cuff tear · Irreparable rotator cuff · Shoulder

Arthroskopische superiore Kapselrekonstruktion mittels humanem dermalem Allograft bei Patienten mit und ohne präoperative Pseudoparalysis

Zusammenfassung

Hintergrund. Irreparable Rotatorenmanschettenrupturen (RMR) können deutliche Schmerzen und Funktionsverlust aufgrund des Verlusts der superioren glenohumeralen Stabilität verursachen. Durch die Rekonstruktion der superioren Kapsel (SCR) wurden Stabilität und Funktion wiedererlangt.

Fragstellung. Ziel der vorliegenden Studie war es, klinische und radiologische Kurzzeitergebnisse nach arthroskopischem SCR-Eingriff mit dermalem Allograft bei irreparabler RMR zu erfassen.

Material und Methoden. In diese Studie wurden 21 Patienten mit einem Durchschnittsalter von 65,9 Jahren eingeschlossen, bei denen eine arthroskopische SCR erfolgte. Zur Graft-Evaluierung wurde eine Untersuchung mittels Magnetresonanztomographie (MRT) vor und 12 Monate nach der Operation durchgeführt.

Das funktionelle und das subjektive Ergebnis wurde mittels Constant Score (CS), American Shoulder and Elbow Surgeons Score (ASES) und den Simple Shoulder Test (SST) vor, 3–6 Monate und 12 Monate nach der Operation evaluiert. Diese Parameter wurden zwischen Patienten mit und ohne präoperative Pseudoparalyse verglichen. Komplikationen wurden dokumentiert.

Ergebnisse. Nach einem Jahr erwies sich in 71,5% der Fälle der Patch in der MRT als intakt, es trat eine signifikante Besserung im CS und dessen Untergruppen Schmerz, Bewegung und Kraft, im ASES und im SST auf (p < 0.01). Eine Pseudoparalyse zeigte sich bei 51,7% aller Patienten präoperativ, diese hatte sich bei allen Patienten mit intaktem Patch nach einem Jahr zurückgebildet. Das klinische Ergebnis war nach einem Jahr in beiden Gruppen vergleichbar. Komplikationen traten in 29% der Fälle auf (1 Infektion, 5 Patchversagen), bei diesen wurde die Indikation zur inversen Prothese gestellt.

Schlussfolgerung. Der arthroskopische SCR mit humanem dermalem Allograft bei Patienten mit irreparabler RMR zeigt eine funktionelle Verbesserung und Schmerzereduktion. Dies wurde sowohl für Patienten mit guter Funktion als auch für Patienten mit glenohumeraler Pseudoparalyse nachgewiesen.

Schlüsselwörter

Pseudoparalyse · Superiore Kapselrekonstruktion · Massenruptur · Irreparable Rotatorenmanschette · Schulter

Statistical analysis

Patient details are presented using descriptive statistics. The distribution of the data was assessed by visual inspection of histograms and the Kolmogorov–Smirnov test. Parametric data are presented as mean ± standard deviation, nonparametric data as median and range in parenthesis, or as numbers and percentages in parenthesis for categorical variables. Paired t tests or nonparametric Wilcoxon signed-rank tests were used to compare parameters between two time points. Independent t tests or nonparametric Mann–Whitney U tests were used to compare continuous variables between two groups. Chi-square or Fisher’s exact tests were performed to analyze categorical variables. All comparative tests were two-tailed and the statistical significance level was set at p < 0.05. Statistical analyses were performed using SPSS Statistics 23 (IBM® Corporation, Armonk, NY, USA).

Results

Demographic data and surgical details of all patients treated with arthroscopic
SCR using a human dermal allograft are presented in Table 1. Arthroscopic surgery prior to SCR included failed supraspinatus tendon reconstructions, latissimus dorsi transfer, debridement and long head of biceps tenotomy. No significant differences regarding demographic data were detected between patients with and without pseudoparalytic conditions (Table 1). Tendon tears involving the musculus supraspinatus, infraspinatus, or subscapularis were detected and reported if reconstruction was needed. Preoperative radiographic evaluation according to the Hamada classification showed eight patients with Hamada Grade 1 and 13 shoulders with Hamada Grade 2. No significant difference in outcome or complications was detected.

The MRI evaluation revealed intact dermal grafts in 15 patients (71.4%; Fig. 1). An MRI-verified rupture of the implanted graft was detected at 6.2 ± 2.8 months after SCR in six out of 21 cases (28.4% total failure rate; location of the rupture in three cases at the humeral side and three at the glenoidal side, Fig. 2). Revision surgery was performed on five patients (23.8%) after a mean of 8.4 ± 4.6 months. Reverse total shoulder arthroplasty was performed in four cases: due to isolated graft failure with progressive cuff tear arthropathy in two cases, and graft failure with additional subscapularis rupture in the other two cases. In one case a low-grade infection (Cutibacterium acnes) resulting in graft failure led to subsequent revision surgeries (debridement and removal of all foreign materials followed by a planned reverse total shoulder arthroplasty). One additional patient with an isolated graft failure is scheduled to undergo reverse total shoulder arthroplasty.

Clinical and subjective evaluations of patients with intact grafts showed significant improvements from baseline to both postoperative time points regarding the CS including its subgroups pain, ADL, ROM and strength, ASES, SST, and the absolute ROMs in forward flexion and abduction (Table 2). Comparisons between the first and second follow-up showed either constant or further improved outcome scores at the 12-month timepoint (Table 2).

Preoperative pseudoparalysis was present in 12 out of 21 cases (57.1%). The majority of patients with SCR failure (four out of six patients, 66.7%) presented with a preoperative pseudoparalysis including three patients with a forward flexion and/or abduction of less than 60°. The 3–6-month and 12-month follow-up revealed reversal of pseudoparalysis in 80 and 100% of patients (revisions not included), respectively (Table 3).

Patients with pseudoparalysis compared with those without pseudoparalytic conditions showed significant baseline differences regarding total Constant scores and ASES (Table 4). The SST was significantly better in the group without than in the group with pseudoparalysis at the 3–6-month follow-up (Table 4). Regarding ROM, only the Constant score was significantly better at baseline and at the 3–6-month follow-up in the group without than in the group with shoulder function of 90° or below, as shown in Fig. 3. From the subgroup analysis, both groups overall showed significant improvement in forward flexion and abduction; for group 1 (forward flexion over 90°), the forward flexion was from 147 to 161°. For group 2, forward flexion of 90° or below showed improvement for group 2a (forward flexion between 60 and 90°) from 74 to 155° and for group 2b (forward flexion below 60°) from 38 to 168°.
### Table 2: All scores before and after arthroscopic SCR with a human dermal allograft

|                       | Baseline | 3–6 months FU1 | 12 months FU2 | p (baseline vs. FU1) | p (baseline vs. FU2) | p (FU1 vs. FU2) |
|-----------------------|----------|----------------|--------------|----------------------|----------------------|-----------------|
| CS Total              | 30.3 ± 15.3 | 61.9 ± 15.1 | 77.3 ± 15.2 | <0.001               | <0.001               | 0.004           |
| CS pain               | 5 (0–5)   | 10 (5–15)    | 15 (10–15)  | 0.001                | 0.001                | 0.025           |
| CS ADL                | 6 (2–17)  | 14 (9–28)    | 18 (8–28)   | 0.001                | 0.001                | 0.029           |
| CS ROM                | 16 (4–38) | 26 (18–40)   | 36 (22–40)  | 0.016                | 0.001                | 0.005           |
| CS strength           | 0 (0–6)   | 10 (2–20)    | 10 (0–25)   | 0.002                | 0.001                | 0.172           |

*ABD* abduction, *ADL* activity of daily living, *CS* Constant score, *FU* follow-up, *ROM* range of motion, *SCR* superior capsular reconstruction, *SST* Simple Shoulder Test

### Table 3: Comparison of scores between patients with and without pseudoparalysis

|                       | Forward flexion >90° | Forward flexion ≤90° | p |
|-----------------------|---------------------|----------------------|---|
| **CS Total**          | (n = 7)             | (n = 8)              |   |
| Baseline              | 40.7 ± 16.2         | 21.1 ± 6.4           | 0.018 |
| 3–6 months            | 68.0 ± 13.8         | 56.5 ± 15.0          | 0.148 |
| 12 months             | 76.0 ± 18.3         | 78.4 ± 13.2          | 0.776 |
| **ASES**              |                     |                      |   |
| Baseline              | 41.0 ± 14.7         | 27.9 ± 6.5           | 0.041 |
| 3–6 months            | 76.7 ± 15.5         | 58.8 ± 21.1          | 0.087 |
| 12 months             | 84.3 ± 18.4         | 80.0 ± 16.1          | 0.638 |
| **SST**               |                     |                      |   |
| Baseline              | 3.3 ± 2.9           | 1.6 ± 1.8            | 0.222 |
| 3–6 months            | 10.3 ± 1.7          | 6.0 ± 2.6            | 0.002 |
| 12 months             | 9.3 ± 3.1           | 7.9 ± 3.5            | 0.428 |

*ASES* American Shoulder and Elbow Surgeons score, *CS* Constant score, *SST* Simple Shoulder Test

### Discussion

The most important finding of the present study was that SCR with an acellular human dermal allograft could be confirmed to be a valid short-term option to restore shoulder function in patients with irreparable supraspinatus tendon tears or with or without repairable subscapularis or infraspinatus and limitation in shoulder function. However, there was a relatively high revision rate in the short term that has to be considered when performing SCR.

After 3–6 months, 80% of patients with pseudoparalysis regained a significant part of their shoulder function. This rate further increased to 100% at the 12-month mark. Burkhart et al. [1] were able to show similar results for SCR in patients with pseudoparalysis and irreparable rotator cuff tear with reproducible reversal of pseudoparalysis in 90% of the cases compared with a significantly lower rate of 44% following partial repairs. Mihata et al. [18] showed significant improvement in range of motion from 54.3 to 146.8° for patients with moderate (<90° forward flexion) and from 36.7 to 150.0° for patients with severe pseudoparalysis (<45° forward flexion). This is comparable to our results using a dermal allograft, which provided improvement of forward flexion from 74 to 155° in moderate and from 38 to 168° in severe pseudoparalysis, respectively. In our non-pseudoparalysis group (>90° forward flexion) forward flexion improved from 147 to 161°, which is similar to results of a recent study reporting 1-year improvements for forward flexion from 120 to 160° [25].

Despite range of motion and function often being the focus of attention, pain relief remains an important factor for improving a patient’s quality of life and performance of common daily activities. The present study found a significant, swift pain reduction in the first 3–6 months after surgery, which is in line with recent publications showing similar pain reduction from the pre- to postoperative period [1, 4, 5, 25]. Furthermore, it has to be confirmed in long-term studies whether SCR prevents or, at a minimum, leads to clinically significant delays in the progression of cuff tear arthropathy. The use of an Inspace balloon remains controversial; recent studies showed an initial significant improvement but higher complication rates after 1 year of follow-up [12, 28]. Clinical studies with longer follow-up and improved implant systems are needed.

Similar to our study, four recent publications about SCR with human dermal allografts reported failure rates between 40 and 60% with mainly humeral side ruptures [4, 11, 15, 25], while recent data on SCR with fascia lata autografts described failure rates between 5 and 10% [3, 18]. Although the literature on SCR with fascia lata is limited, the healing rate of autografts seems to be far better. The same is known for anterior cruciate ligament reconstruction where the use of allografts is known to be safe, but can be associated with a significantly higher failure
and reoperation rate and postoperative stability \[2, 9, 31\]. Allograft incorporation to host tissue, which can be up to 1 year, takes much longer compared with autografts \[5\]. However, when using autografts, donor site morbidity, especially when taking a long fascia lata graft, needs to be considered. In fact, Azevedo et al. \[3\] reported that at the 2-year follow-up, 57% of patients were still bothered by their harvested thigh and 76% noticed donor site changes.

However, Plachet al. \[26, 27\] in their case report showed that there is ingrowth of vessels and expression of growth factors in the acellular human dermal allograft patch after 6 months, which led them to the conclusion that healing of the graft to the native tissue may be facilitated. However, in their histologic samples they only analyzed soft tissue and not the patch–bone interface, which is crucial for the construct to survive. However, Plachet al. \[26, 27\] in their case report showed that there is ingrowth of vessels and expression of growth factors in the acellular human dermal allograft patch after 6 months, which led them to the conclusion that healing of the graft to the native tissue may be facilitated. However, in their histologic samples they only analyzed soft tissue and not the patch–bone interface, which is crucial for the construct to survive.

Another detail complicating comparisons between autograft and allograft SCR is that to date most studies using allografts started with a preoperatively better shoulder function than the studies using autografts \[6, 27\]. Further definitions of the criteria for the indication for SCR still need to be developed to help decide who will benefit the most from the procedure. Moreover, irreparability of rotator cuff tears is very much surgeon dependent. In our 45 patients who had been initially scheduled for SCR based on history, examination, and preoperative MRI, more than half turned out to have repairable rotator cuff tears with extensive release techniques.

As shown in a current publica-
tion, cuff tear arthropathy at time of surgery remains a limiting factor for expected outcomes. Denard et al. reported differences in success rates of 75% for Hamada Grades 1–2 and 44% for Hamada Grades 3–4. However, our finding together with those from recent publications present promising short- to mid-term results after SCR in patients with irreparable posterosuperior rotator cuff tear under appearance of severe loss in range of motion. Long-term results on consistent regain of function and

**Fig. 3** a Illustration of the mean Constant Score subgroups a pain, b activities of daily living (ADL), c range of motion (ROM), and d strength at baseline and at the 3–6- and 12-month follow-up (FU). Asterisks show significant \((p < 0.05)\) differences between patients with and without preoperative pseudoparalytic conditions.
The primary limitations of this study are the short follow-up period and the low number of enrolled patients. Furthermore, comparisons between other joint-preserving techniques such as debridement, partial rotator cuff repair, or tendon transfers and superior capsular reconstruction should be performed in the future.

### Practical conclusion

- Superior capsular reconstruction (SCR) is a promising option for the treatment of irreparable rotator cuff tears in patients with no or mild osteoarthritis.
- The technique reproducibly provided early reduction of pain and restoration of shoulder function in patients with and without pseudoparalysis.
- However, the short-term revision rate was relatively high, which has to be considered when performing SCR.

### Corresponding address

Dr. Philipp R. Heuberer, MD  
Health Pi  
Wollzeile 1–3, 1010 Vienna, Austria  
philipp@heuberer.at

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**Author Contribution.** All authors made substantive intellectual contributions to the study. The authors’ responsibilities were as follows: P.R. Heuberer, L. Pauzenberger, and W. Anderl initiated the study regarding conception and design; M. Eigenschink and R.C. Ostermann collected data and performed the literature search. B. Laky and M. Eigenschink performed statistical analysis, P.R. Heuberer and W. Anderl supervised the conduction of the study; all authors were actively involved in drafting the manuscript or revising it critically for important intellectual content; and have given final approval of the version to be published.

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### Table 5  Study details of reported arthroscopic superior capsular reconstruction with auto- and allografts

| Publication | Graft type | N  | FU (in years) | Clinical outcome (pre- to postoperative) | ROM (pre- to postoperative) | Healing rate [%] |
|-------------|------------|----|---------------|----------------------------------------|------------------------------|------------------|
| Mihata et al. [18] Arthroscopy 2013 | Fascia lata | 24 | 2–4 | ASES: 24 → 93 | 84° → 148° | 83 |
| Mihata et al. [17] AmJSM 2018 | Fascia lata | 88 | 3–9 | ASES: (44, 29, 20) → (97, 92, 92) | (143°, 54°, 37°) → (164°, 147°, 150°) | 92 |
| Azevedo et al. [3] OJSM 2018 | Fascia lata | 22 | 2 | CS: 18 → 65 | 75° → 144° | 91 |
| Pennington et al. [24] Arthroscopy 2018 | Allograft | 86 | 1 | ASES: 52 → 82 | 120° → 160° | 95 |
| Denard et al. [4] Arthroscopy 2018 | Allograft | 59 | 1 | ASES: 44 → 78 | 130° → 150° | 45 (75) |
| Burkhardt and Hartzler [1] Arthroscopy 2018 | Allograft | 10 | 1 | ASES: 52 → 89 | 27° → 159° | 70 |
| Lee and Min [15] KSSTA 2018 | Allograft | 36 | 2 | ASES: 50 → 84 CS: 56 → 83 | 107° + 158° → 156° + 135° | 64 |
| Hirahara et al. [10] AOJ2019 | Allograft | 9 | 2 | ASES: 42 → 87 | x | 80 (4/5) |
| Present study | Allograft | 21 | 1 | ASES: 34 (41, 28) → 82 (84, 80) CS: 30 (41, 21) → 77 (76, 78) | FF: 107° (147°, 72°) → 161° (161°, 161°) ABD: 100° (144°, 63°) → 156° (156, 156°) | 71 |

**Table** 5. Study details of reported arthroscopic superior capsular reconstruction with auto- and allografts

**ABD**: abduction, **ASES**: American Shoulder and Elbow Surgeons score **CS**: Constant score, **FF**: forward flexion, **FU**: follow-up, **ROM**: range of motion
Compliance with ethical guidelines

Conflict of interest. W. Anderl, R.C. Ostermann, and P.R. Heuberer receive personal fees from Arthrex, Inc. The company had no influence on the study design, data collection, interpretation of the results, or the final manuscript. The remaining authors (M. Eigenschink, L. Pausenberger, B. Laky) declare that they do not have any conflicts of interest pertinent to this study.

The study was approved by the institutional review board (201609_EK13) and all investigations were accomplished regarding national law and the declaration of Helsinki 1975 (revised version). Informed consent was obtained from all patients.

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References

1. Burkhart SS, Hartzler RU (2019) Superior capsular reconstruction reverses profound pseudoparalysis in patients with irreparable rotator cuff tears and minimal or no Glenohumeral arthritis. Arthroscopy 35:22–28
2. Condello V, Zdanowicz U, Di Matteo B et al (2019) Allograft tendons are a safe and effective option for revision ACL reconstruction: a clinical review. Knee Surg Sports Traumatol Arthrosc 27(6):1771–1781. https://doi.org/10.1007/s00167-018-5147-4
3. De Campos Azevedo CJ, Angela A, Vinga S (2018) Arthroscopic superior capsular reconstruction with a minimally invasive harvested fascia Lata Autograft produces good clinical results. Orthop J Sports Med:2325967118808242
4. Denard PJ, Brady PC, Adams CR et al (2018) Preliminary results of Arthroscopic superior capsular reconstruction with dermal Allograft. Arthroscopy 34:93–99
5. Eagan MJ, Mccallister DR (2009) Biology of allograft incorporation. Clin Sports Med 28:203–214 [vi][vii]
6. Ek ET, Neukom L, Catanzaro S et al (2013) Reverse total shoulder arthroplasty for massive irreparable rotator cuff tears in patients younger than 65 years old: results after five to fifteen years. J Shoulder Elbow Surg 22:1199–1208
7. El-Azab HM, Rott O, Irlenbusch U (2015) Long-term follow-up after Iatissimus dorsi transfer for irreparable posterosuperior rotator cuff tears. J Bone Joint Surg Am 97:462–469
8. Familiari F, Rojas J, Nedim Doral M et al (2018) Reverse total shoulder arthroplasty. EFT Open Rev 3:58–69
9. Grassi A, Niti M, Moulton SG et al (2017) Does the type of graft affect the outcome of revision anterior cruciate ligament reconstruction? a meta-analysis of 32 studies. Bone Joint J.199:B714–723
10. Hamada KFH, Mikasa M, Kobayashi Y (1990) Roentgenographic findings in massive rotator cuff tears A long-term observation. Clin Orthop Relat Res 254:92–96
11. Hiranaka AM, Andersen WJ, Panero AJ (2017) Superior capsular reconstruction: clinical outcomes after minimum 2-year follow-up. Am J Orthop (Belle Mead, NJ) 46:266–278
12. Holschen M, Brand F, Agneskirchner J (2017) Subacromial spacer implantation for massive rotator cuff tears: Clinical outcome of arthroscopically treated patients. Obere Extremittät 12:38–45
13. Ishihara Y, Mihata T, Tamboli M et al (2014) Role of the superior shoulder capsule in passive stability of the glenohumeral joint. J Shoulder Elbow Surg 23:642–648
14. Klinger HM, Spahn G, Baums MH et al (2005) Arthroscopic debridement of irreparable massive rotator cuff tears—a comparison of debridement alone and combined procedure with biceps tenotomy. Acta Chir Belg 105:297–301
15. Lee SJ, Min YK (2018) Can inadequate acromio-humeral distance improvement and poor posterior remnant tissue be the predictive factors of re-tear? Preliminary outcomes of arthroscopic superior capsular reconstruction. Knee Surg Sports Traumatol Arthrosc 26:2205–2213
16. Ling HY, Angeles JG, Horodyski MB (2009) Biomechanics of Iatissimus dorsi transfer for irreparable posterosuperior rotator cuff tears. Clin Biomech (Bristol, Avon) 24:261–266
17. Merolla G, Porcellini G (2014) Reverse shoulder Arthroplasty in patients aged sixty years old or younger: are we really doing the best? Transl Med UniSa 9:66–67
18. Mihata T, Lee TQ, Hasegawa A et al (2018) Arthroscopic superior capsule reconstruction can eliminate pseudoparalysis in patients with irreparable rotator cuff tears. Am J Sports Med 46:2707–2716
19. Mihata T, Lee TQ, Watanabe C et al (2013) Clinical results of arthroscopic superior capsule reconstruction for irreparable rotator cuff tears. Arthroscopy 29:459–470
20. Mihata T, Mccgarry MH, Kahn T et al (2016) Biomechanical effect of thickness and tension of fascia Lata graft on Glenohumeral stability for superior capsule reconstruction in irreparable Supraspinatus tears. Arthroscopy 32:418–426
21. Mihata T, Mccgarry MH, Kahn T et al (2016) Biomechanical role of capsular continuity in superior capsule reconstruction for irreparable tears of the Supraspinatus tendon. Am J Sports Med 44:1423–1430
22. Mihata T, Mccgarry MH, Pirollo JM et al (2012) Superior capsule reconstruction to restore superior stability in irreparable rotator cuff tears: a biomechanical cadaveric study. Am J Sports Med 40:2249–2255
23. Muh SJ, Streit JJ, Wanner JP et al (2013) Early follow-up of reverse total shoulder arthroplasty in patients sixty years of age or younger. J Bone Joint Surg Am 95:1877–1883
24. Patte D (1990) Classification of rotator cuff lesions. Clin Orthop Relat Res. https://doi.org/10.1097/00003086-199005000-00012
25. Pennington WT, Bartz BA, Pauli JM et al (2018) Arthroscopic superior capsular reconstruction with Acellular dermal Allograft for the treatment of massive irreparable rotator cuff tears: short-term clinical outcomes and the radiographic parameter of superior capsular distance. Arthroscopy 34:1764–1773
26. Pielach F, Klatte-Schulz F, Minkus M et al (2018) Biological allograft healing after superior capsule reconstruction. J Shoulder Elbow Surg 27:e387–e392
27. Plachef F, Minkus M, Scheibel M (2018) Superior Kapselrekonstruktion: Indikation und klinische Resultate. Obere Extremittät 13:237–245
28. Singh JA, Sperling J, Buchbinder R et al (2011) Surgery for shoulder osteoarthritis: a Cochrane systematic review. J Rheumatol 38:598–605
29. Sugaya HMK, Matsuki K, Morishii J (2007) Repair integrity and functional outcome after arthroscopic double-row rotator cuff repair. A prospective outcome study. J Bone Joint Surg Am 89:953–960
30. Tokish JM, Alexander TC, Kissenberth MJ et al (2017) Pseudoparalysis: a systematic review of term definitions, treatment approaches, and outcomes of management techniques. J Shoulder Elbow Surg 26:e177–e187
31. Wang HD, Zhang H, Wang TR et al (2018) Comparison of clinical outcomes after anterior cruciate ligament reconstruction with hamstring tendon autograft versus soft-tissue allograft: a meta-analysis of randomised controlled trials. Int J Surg 56:174–183