



Effect of complications on outcomes after revision reverse total shoulder arthroplasty

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Background: Revision reverse total shoulder arthroplasty (RTSA) reliably improves shoulder pain and function in patients with failed shoulder arthroplasty, although it can lead to significant postoperative complications. The purpose of this study was to determine the effect of postoperative complications on shoulder pain and function after revision RTSA.

Methods: We evaluated 36 patients at an average of 4.3 years (range, 2–8.6 years) after revision of a shoulder arthroplasty to RTSA. Of these patients, 9 had a failed anatomic total shoulder arthroplasty, 23 had a failed hemiarthroplasty, and 4 had a failed RTSA. The American Shoulder and Elbow Surgeons (ASES) score and visual analog scale (VAS) pain score were evaluated postoperatively, and patients with and without postoperative complications were compared.

Results: The final ASES score and VAS pain score were 61 ± 23 and 2.4 ± 2.3 , respectively. A major postoperative complication occurred in 7 patients (19%) (infection in 3, hematoma in 1, instability in 1, and acromial and/or scapular spine fracture in 2). Further surgical treatment was required in 5 patients (14%) (irrigation and débridement and component exchange for infection in 3, irrigation and débridement for hematoma in 1, and open reduction—internal fixation of scapular spine fracture in 1). On comparison of clinical outcomes between patients with and patients without complications, the ASES score and VAS pain score were significantly worse in patients with complications vs. those without them (ASES score, 43 ± 24 vs. 66 ± 21 [$P = .04$]; VAS pain score, 4.3 ± 2 vs. 2 ± 2.2 [$P = .03$]).

Conclusion: Revision RTSA resulted in postoperative pain and shoulder function comparable to primary RTSA reported in the literature, although postoperative complications led to clinically significant declines in function and increases in pain.

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As the incidence of shoulder arthroplasty has increased,²⁸ especially in younger patients,^{33,39} the burden of revision shoulder arthroplasty has increased.^{6,50} Historically, revision total shoulder arthroplasty (TSA) was associated with unacceptably high failure rates because of difficulty reconstructing glenoid bone defects,^{17,21,24} difficulty achieving stable fixation of the glenoid component,¹¹ and a high prevalence of rotator cuff failure.³⁷ The expectation after revision TSA in the past was the achievement of limited goals with modest levels of pain relief or functional improvement. Revision reverse total shoulder arthroplasty (RTSA) eliminates many problems associated with revision TSA. Glenoid

bone grafting can be very successful in the setting of RTSA,^{8,25,50,56} baseplate loosening is uncommon,^{22,30,43} and RTSA does not rely on an intact rotator cuff.^{10,19} Unfortunately, revision RTSA comes at the cost of other unique postoperative complications including impingement resulting in instability and acromial or scapular spine fractures, which may affect final functional outcomes. Varying results of revision RTSA have been reported in several studies, with improvements in pain and function, although complication rates have reached as high as 50%^{4,7,18,22,23,27,31,35,36,48,50,53} (Table 1).

Saltzman et al³⁸ previously reported on the complication rates of primary vs. revision RTSA and TSA and determined that overall, minor, surgical, intraoperative, perioperative, and postoperative complications were greater after revision RTSA than after primary RTSA. Very few studies have examined the influence of postoperative complications in revision RTSA on functional outcomes. Holcomb et al²² reported the results of revision RTSA owing to baseplate failure in RTSA and found no difference in outcomes comparing pre-failure vs. post-revision American Shoulder and

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Elbow Surgeons (ASES) scores, supporting that the outcomes of primary vs. revision RTSA show no difference. Shields and Wiater⁴¹ reported on 36 revision RTSAs and noted significantly worse Subjective Shoulder Values in the revision RTSA group when patients sustained either a postoperative complication or required revision surgery compared with patients who did not require revision surgery or sustain a postoperative complication.

The purpose of this study was to describe the short-term clinical and radiographic results of revision to RTSA and determine whether patients sustaining a postoperative complication had worse functional results than those without a complication. We hypothesized that revision to RTSA would result in significant improvements in range of motion and functional outcomes and that patients sustaining a postoperative complication would have significantly worse shoulder pain and function.

Materials and methods

This was a retrospective study. The operative log of the senior author (R.Z.T.) was reviewed. We queried the log for patients who underwent revision from a humeral hemiarthroplasty (HA), anatomic TSA, or RTSA to an RTSA from May 2008 to May 2015. Patient who underwent conversion from an HA, anatomic TSA, or RTSA to an antibiotic spacer, anatomic TSA, or HA were excluded. We also excluded patients who underwent primary RTSA and patients with <2 years' potential follow-up. RTSA was indicated for patients with persistent pain, limited function, and restriction of motion in those with a failed arthroplasty. In the setting of a failed HA, the indications for conversion to RTSA included a painful arthroplasty with or without rotator cuff insufficiency with or without glenoid or humeral bone deficiency. No HA was converted to an anatomic TSA during the study period. In the setting of a failed anatomic TSA, the indications for conversion to RTSA included a

loosened glenoid implant not amendable to revision anatomic glenoid component placement or rotator cuff failure. In the setting of a failed RTSA, indications for conversion to another RTSA included instability or implant loosening amendable to revision with or without bone grafting. Eligible patients were then contacted and returned for evaluation including functional outcome questionnaires, physical examinations, and radiographs or provided functional outcome data by telephone.

Operative protocol

In all cases, a deltopectoral approach was used. Both the Trabecular Metal RTSA (Zimmer, Warsaw, IN, USA) and Aequalis Reversed Shoulder Arthroplasty (Wright Medical Group, Bloomington, MN, USA) systems were used. Although initial attempts were made to free the humeral stem using a burr and thin osteotomes, when this method was not successful an osteotomy or window of the humeral shaft was performed and was fixated with cerclage wires as previously described.⁷ Both long (>130 mm) and standard-length (130 mm) stemmed implants were used depending on proximal humeral bone loss and the requirement for a humeral osteotomy or window. In cases of severe glenoid erosion, allograft or iliac crest autograft reconstruction of the glenoid defect was performed. Glenoid grafting was performed in cases in which severe cavitory defect limited the ability to implant a baseplate in a stable fashion or severe erosion limited the baseplate to be positioned in a neutral position or with inferior tilt. We used 36-, 40-, and 42-mm glenospheres depending on the implant system, as well as the size of the patient. Standard or constrained polyethylene liners were used at the discretion of the surgeon on the basis of overall stability of the shoulder during trialing. The subscapularis was not repaired in any case. Proximal humeral allograft was used in some cases at the discretion of the surgeon for proximal humeral

Table 1
Results of revision to reverse total shoulder arthroplasty in prior series

Authors	Year	Patients, n	Follow-up, mo	AFE, °	VAS pain score	ASES score	Constant score	SST score	Cx, %
Alentorn-Geli et al ¹	2017	31	28	109	1	NA	NA	NA	21
Black et al ⁴	2015	16	59	NA	1.7	67	NA	5.3	56
Black et al ⁵	2014	36	55	NA	1.4	70	NA	5.9	28
Boileau et al ⁷	2013	37	34	111	NA	NA	47	NA	30
Castagna et al ¹²	2013	36	32	120	NA	NA	48	NA	0
Chacon et al ¹³	2009	25	30	82	NA	70	NA	4.5	16
Cox et al ¹⁵	2019	73	68	75	NA	51	NA	3.5	19
Flury et al ¹⁸	2011	21	46	97	3	NA	56	NA	43
Hernandez et al ²⁰	2017	70	36	112	NA	68	NA	7	NA
Holcomb et al ²²	2009	14	33	118	NA	70	NA	4.5	21
Holschen et al ²³	2017								
GHOA cohort		23	24	126	1.5	59	67	NA	9
Fracture cohort		21	24	115	2.2	71	73	NA	24
Kany et al ²⁶	2015	29	28	124	2	NA	60	8	0
Kelly et al ²⁷	2012	28	34	106	NA	72	49	NA	50
Melis et al ³¹	2012	37	47	121	NA	NA	55	NA	30
Merolla et al ³²	2018	157	49	NA	2.2	60	NA	6	NA
Ortmaier et al ³⁵	2013	50	51	98	1	NA	49	5.6	24
Patel et al ³⁶	2012	28	41	108	2.6	66	NA	7.6	11
Sheth et al ⁴⁰	2019	110	57	NA	2.9	63	NA	NA	20
Shields and Wiater ⁴¹	2019	35	50	NA	2.4	68	NA	NA	31
Stephens et al ⁴³	2016	58	24	97	3.5	53	NA	4.4	18
Stephens et al ⁴⁵	2015	16	36	100	2.6	67	NA	5.3	31
Valenti et al ⁴⁸	2014	30	36	108	NA	NA	52	NA	27
Wagner et al ⁵⁰	2015	143	37	NA	NA	66	NA	6	18
Wagner et al ⁵²	2017	39	36	121	NA	68	NA	6.8	15
Wagner et al ⁵¹	2017	38	44	108	NA	61	NA	5	26
Walker et al ⁵³	2012	22	40	130	NA	68	NA	5	23
Weighted mean or total	NA	1223	NA	106 ± 15	3 ± 2	63 ± 6	54 ± 8	5.7 ± 1.1	22

AFE, active forward elevation; VAS, visual analog scale; ASES, American Shoulder and Elbow Surgeons; SST, Simple Shoulder Test score; Cx, complications; NA, not available; GHOA, glenohumeral osteoarthritis.

Data are presented as mean values unless otherwise indicated.

Table II
Clinical results of patients with complications or need for reoperation vs. patients without complications or need for reoperation

	Patients, n	Final ASES score	Final VAS pain score
Complications			
Yes	7	43 ± 24	4.3 ± 2
No	29	66 ± 21	2 ± 2.2
P value		.04	.03
Reoperations			
Yes	5	45 ± 28	3.8 ± 1.9
No	31	64 ± 22	2.2 ± 2.2
P value		.22	.21

ASES, American Shoulder and Elbow Surgeons; VAS, visual analog scale. The Student *t* test was used to compare outcome scores; *P* < .05 was considered statistically significant.

bone loss in the setting of tuberosity resorption using the step-cut technique described by Chacon et al.¹³

Clinical data collection

For each patient, the following data were collected based on the preoperative documentation: age, operative side, sex, duration of preoperative symptoms, active forward elevation (AFE) of the shoulder, and whether the patient had an active workers' compensation claim. In addition, for each patient, the following data were collected based on the intraoperative documentation: diagnosis, procedure, implant company, whether a glenoid bone graft was required and the source of this graft, whether an allograft was required for proximal humeral bone deficiency, whether a constrained polyethylene liner was used, glenosphere size, glenoid baseplate peg length, whether a humeral osteotomy was required, whether a humeral window was required, length of the humeral stem (standard or long stem), and whether any intraoperative complications or fractures were noted. For each patient, the following data were collected at final follow-up: whether any major postoperative complication occurred (fracture, instability, and infection or hematoma requiring surgery), whether the patient underwent revision, AFE, visual analog scale (VAS) pain score, ASES score, and length of follow-up.

Radiographic data collection

Preoperative and most recent final follow-up radiographs, including anteroposterior, Grashey anteroposterior, scapular-Y lateral, and axillary lateral views, were independently evaluated by an attending surgeon (P.N.C.) who had fellowship training in shoulder and elbow surgery but did not perform the index procedures. Preoperative humeral bone deficiency was classified using the PHAROS (Proximal Humeral Arthroplasty Revision Osseous Insufficiency) classification system.¹⁴ Postoperative radiographs were evaluated for incorporation or resorption of the bone grafts, migration or subsidence of the baseplate, humeral stem loosening, scapular spine or acromial fracture, proximal humeral osteolysis, and scapular notching. Scapular notching was graded using the Nerot-Sirveaux system.⁴²

Statistical analysis

Descriptive statistics were calculated for preoperative, intraoperative, and postoperative variables. Preoperative AFE and postoperative AFE were compared using paired Student *t* tests. Postoperative outcome scores were compared between patients with and patients without complications using Student *t* tests. A power analysis was not performed because the procedure is an

uncommon procedure that was examined with a retrospective design. All available subjects were included.

Results

In total, 56 patients met the inclusion and exclusion criteria, of whom 8 died, leaving a total of 48 patients available for review. Of these 48 patients, 12 were lost to follow-up, resulting in a total of 36 patients being reviewed and a 75% rate of follow-up. The length of follow-up was 4.3 years (range, 2–8.6 years). The right side was affected in 66% of patients. Female patients comprised 86% of the cohort, and the mean age (±standard deviation) was 69 ± 11.3 years. One patient had a prior workers' compensation claim. Preoperatively, patients showed significant restriction of motion, with AFE of only 55° ± 46.9°. Of the patients, 9 had a failed anatomic TSA (rotator cuff tear in 6, loosened glenoid component in 2, and periprosthetic fracture in 1), 23 had a failed HA (antibiotic spacer in 4 and rotator cuff failure or tuberosity nonunion and/or glenoid erosion in 19), and 4 had a failed RTSA (infection in 1, instability in 1, and loosened glenoid component in 2).

Preoperatively, 4 patients (11%) had no humeral bone loss. In contrast, 23 patients (64%) had type 1 epiphyseal humeral bone loss (15 with cortical thinning [type 1A], 3 with loss of the calcar [type 1AC], 4 with loss of the greater tuberosity [type 1AG], and 1 with loss of the greater tuberosity and calcar [type 1ACG]); 6 patients (17%) had type 2 metaphyseal bone loss (4 with cortical thinning [type 2A] and 2 with cortical loss [type 2B]); and 3 patients (8%) had type 3 diaphyseal bone loss (2 with cortical thinning [type 3A] and 1 with cortical loss [type 3B]).

Intraoperatively, a Trabecular Metal RTSA was used in 14 cases (39%) whereas an Aequalis Reversed Shoulder Arthroplasty was used in 22 cases (61%). A constrained polyethylene liner was used in 5 patients (14%). The humeral stem was not revised in 5 cases, whereas a long-stem implant (>130 mm) was used in 7 cases and a standard stem (130 mm) was used in 24 cases. A 36-mm glenosphere was used in all cases. In 8 cases, a 25-mm extended-length baseplate peg was used, whereas the remaining baseplates had a 15-mm-long central post. An episiotomy was required in 3 patients to remove the stem, and a humeral window was required in 2. Four patients underwent glenoid bone grafting with femoral head allograft for a large structural defect, and one underwent iliac crest autograft augmentation for a large glenoid defect. Five patients underwent proximal humeral allografting for large proximal humeral bone defects. Two intraoperative complications occurred; both were greater tuberosity fractures repaired with cerclage suture at the time of revision.

On analysis of final radiographs obtained at an average of 2.6 ± 2 years postoperatively, 31 of 36 patients (86%) had well-fixed components without osteolysis or loosening, 1 patient had a displaced scapular fracture that precluded full determination of whether the glenoid component was well fixed, 1 patient had a loose humeral component, and 3 patients had proximal humeral osteolysis but without loosening. By use of the Nerot-Sirveaux system, 20 patients (56%) had no notching, 9 (26%) had grade 1 notching, 1 (3%) had grade 2 notching, 1 (3%) had grade 3 notching, and 4 (11%) had grade 4 notching; in 1 patient, notching could not be evaluated because of a displaced scapular fracture. Of the 10 patients with humeral or glenoid grafts, 9 showed healing whereas 1 could not be evaluated because of a displaced scapular fracture.

Final postoperative AFE was 117° ± 43°, which improved by 62° ± 51° from preoperatively (*P* < .0001). The final ASES score and VAS pain score were 61 ± 23 and 2.4 ± 2.3, respectively. A major postoperative complication occurred in 7 patients (19%) (infection in 3, hematoma in 1, instability in 1, and acromial and/or scapular spine fracture in 2). Further surgical treatment was required in 5 patients

(14%) (3 underwent irrigation and débridement and component exchange for infection, 1 underwent irrigation and débridement for hematoma, and 1 underwent open reduction–internal fixation of a scapular spine fracture that went on to achieve union although the final ASES and VAS pain scores were poor, at 25 and 5, respectively). The humeral stem and the baseplates were maintained in all patients until final follow-up. On comparison of clinical outcomes between patients with and patients without complications, the ASES score and VAS pain score at final follow-up were significantly worse in patients with complications vs. those without them (ASES score, $P = .04$; VAS pain score, $P = .03$) (Table II). The ASES score and VAS pain score were also worse in those patients requiring revision surgery vs. those not requiring revision, although the differences did not achieve statistical significance (ASES score, $P = .22$; VAS pain score, $P = .21$).

Discussion

Revision RTSA has been shown to reliably improve pain and function in the treatment of failed anatomic and reverse shoulder arthroplasty. Despite high rates of clinical success, complication rates are high, ranging up to 50%. Our data are comparable to those of other studies reporting clinical results, with an ASES score averaging 61 and VAS pain score averaging 2.3 with final average AFE of 117°. Of the patients, 19% sustained a postoperative complication and 14% required further surgery. We have also shown a 90% rate of radiographic healing of bulk humeral and glenoid allograft reconstructions, with only 1 baseplate and 1 humeral implant showing loosening. On the basis of our data, avoidance of hematoma, fracture, infection, and instability after revision RTSA is critical to success as these

complications will result in a clinically significant decline in shoulder function.

Multiple prior studies have examined the results of RTSA as revision from a prior HA, TSA, or RTSA.^{4,5,13,23,26,27,29,31,34–36,44,48–50,53,54} Nearly all prior reports have suggested that revision RTSA does decrease patient pain and improve patient-reported function.^{4,5,13,22,23,26,27,29,31,34–36,43,44,48–50,54} However, numerous reports have also noted that final postoperative outcomes after revision RTSA are inferior to those after primary RTSA.^{2,5,9,54} In addition, several reports have noted concerning complication rates as high as 50%–69%.^{4,18,23,27,39,54} Within these reports, predictors of complications and inferior outcomes included a high body mass index,³⁸ medical comorbidities,⁵⁷ glenoid bone grafting,⁵⁰ and proximal humeral bone loss.⁴⁴ Overall, our results are comparable to the findings in the prior literature regarding shoulder function, pain, and complication rates. Unlike prior studies in which patients were not stratified by complications, our study has shown that revision RTSA can lead to similar outcomes to primary RTSA as long as complications are avoided. Only Shields and Wiater⁴¹ determined that the development of a postoperative complication negatively influenced outcomes. Our data support the finding that if a major postoperative complication occurs (Fig. 1), pain and functional outcome will decline, and the inferior results are clinically significant as the differences exceed the minimal clinically important differences previously reported for the ASES score and VAS pain score.⁴⁷

Glenoid and humeral bone graft is common with revision RTSA, although the ideal glenoid or humeral grafting technique remains unclear. Various authors have reported on structural glenoid bone grafting,^{16,46,55} bulk proximal humeral allografting,¹³ femoral neck allografting for glenoid defects,³ and a BIO-RSA technique for

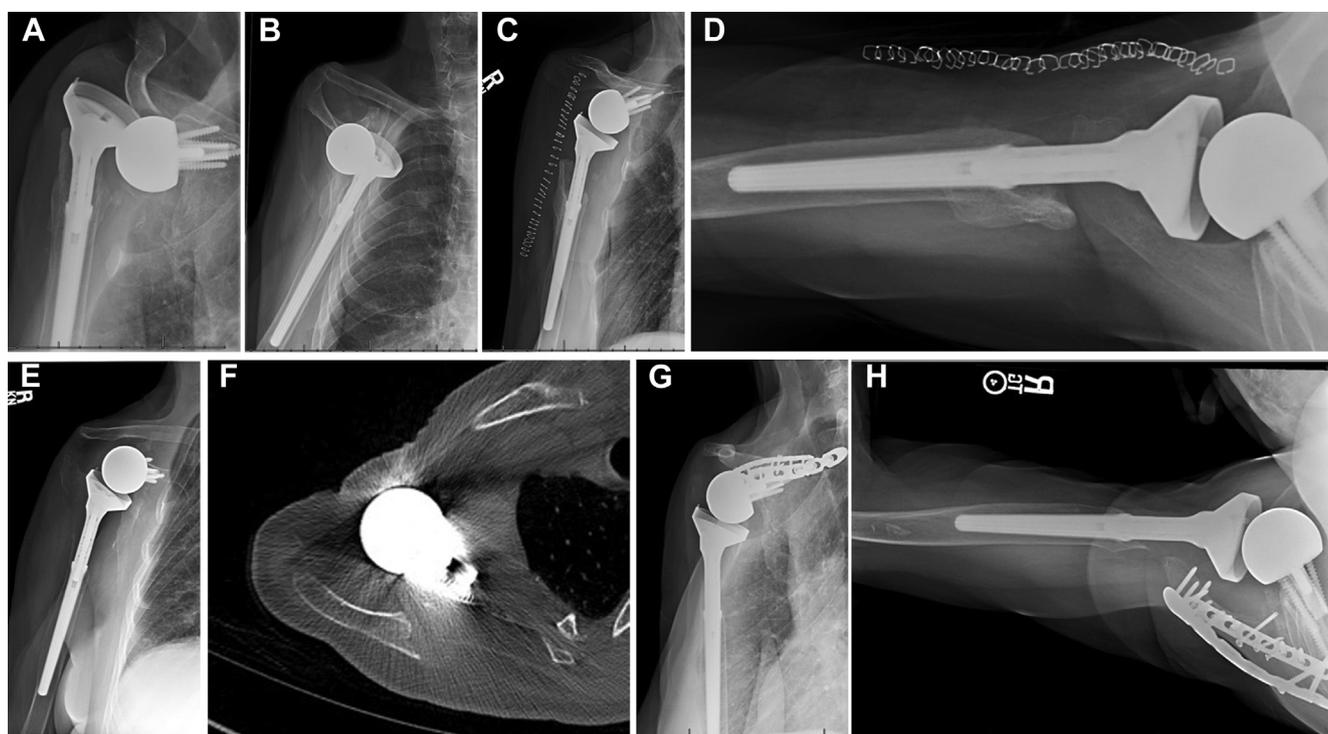


Figure 1 Anteroposterior (A) and axillary lateral (B) radiographs showing a 70-year-old female patient who underwent revision of a recurrently unstable reverse total shoulder arthroplasty with cerclage of the glenosphere to the humerus and 15 mm of lengthening on the humeral side. Immediately postoperatively, anteroposterior (C) and axillary lateral (D) radiographs showed a good outcome. However, a displaced scapular spine fracture occurred at 1.5 years postoperatively; an anteroposterior radiograph (E) and axillary computed tomography slice (F) demonstrate displacement. This progressed to a nonunion, which was painful and limited elevation; thus, the patient underwent open reduction–internal reduction (G, H). Her function remained limited, with final active forward elevation of 30°, and a final American Shoulder and Elbow Surgeons score of 25.

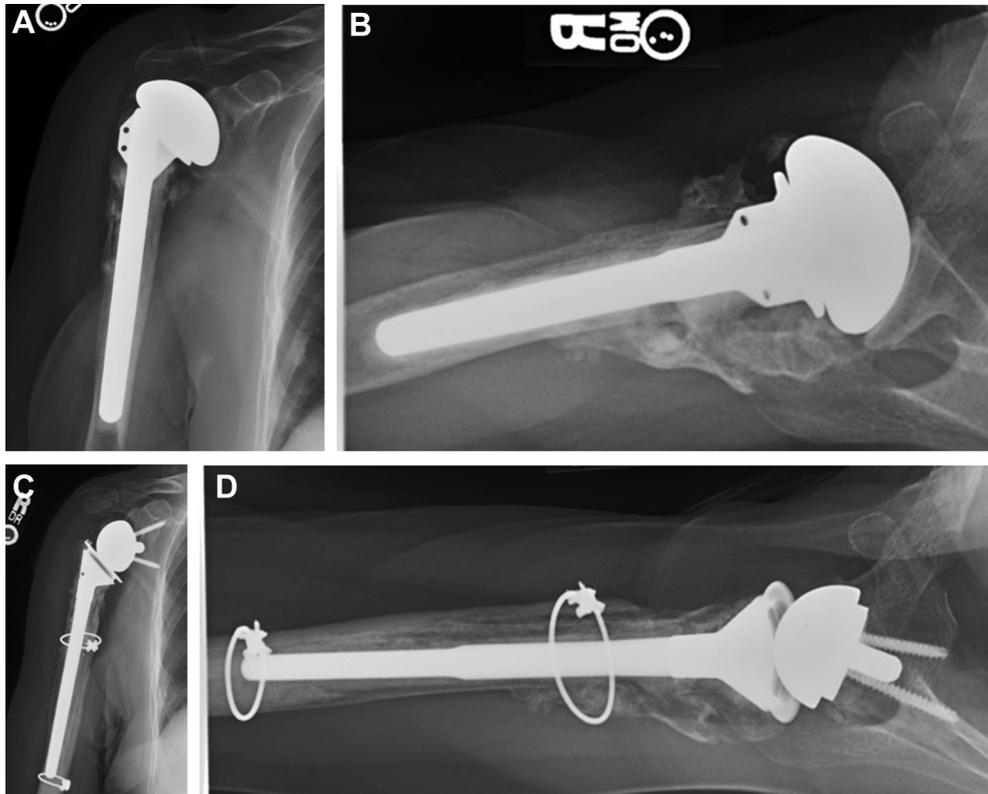


Figure 2 Anteroposterior (A) and axillary lateral (B) radiographs showing an 87-year-old female patient who underwent revision of a painful cemented hemiarthroplasty performed for a proximal humeral fracture with malunion of the greater tuberosity with a proximal humeral allograft and femoral strut allograft. (C, D) Stable implants and graft incorporation were observed at 4 years postoperatively. The patient did not have any complications and was satisfied with her final active forward elevation.

glenoid defects.⁶ Wagner et al⁵⁰ examined a series of 143 revision RTSAs with a minimum follow-up period of 2 years, of which 41 received glenoid bone grafts. Glenoid bone grafting was associated with decreased prosthesis survival at 2 and 5 years' follow-up. Within the series, allograft vs. autograft and corticocancellous vs. structural bone grafting did not influence survival, although the study may have been underpowered for these comparisons.^{46,50} Bulk humeral allografting for proximal humeral defects was evaluated by Chacon et al,¹³ who reported 84% metaphyseal healing and 76% diaphyseal healing with ASES scores averaging 69 at an average of 30 months. Implant survival in our series was not affected by the presence or absence of glenoid or humeral bone grafting using allograft (Fig. 2) as all implants were in place at final follow-up and all bone grafts that could be evaluated by postoperative imaging were healed.

The strengths of this study include a high rate of follow-up; the inclusion of validated, patient-related outcome scores; the comparison of outcomes in patients with and without postoperative complications; and the inclusion of radiographic outcomes. Our study also has several limitations: This was a single-center, retrospective review of a small sample with short-term follow-up and no control group. Given the relative rarity of revision RTSA compared with primary RTSA, larger sample sizes might be difficult to obtain, and nonsignificant differences in analyses could be due to a type II error.

Conclusion

Revision RTSA resulted in postoperative pain and shoulder function comparable to primary RTSA reported in the literature, although postoperative complications led to clinically significant declines in function and increases in pain.

Disclaimer

Robert Z. Tashjian is a paid consultant for Zimmer/Biomet, Wright Medical, and Mitek; owns stock in Conexions, Intrafuse, Genesis, and Kator; receives intellectual property royalties from Wright Medical, Shoulder Innovations, and Zimmer/Biomet; receives publishing royalties from the *Journal of Bone and Joint Surgery* and Springer; and serves on the editorial boards of the *Journal of the American Academy of Orthopaedic Surgeons* and *Shoulder & Elbow*.

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