Radial Shortening Osteotomy for Symptomatic Kienböck's Disease: Complications and Long-Term **Patient-Reported Outcome**

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Abstract

Objective To determine the rate of salvage procedures and any other unplanned reoperations in patients with symptomatic Kienböck's disease who were treated with radial shortening osteotomy. In addition, we studied patient-reported outcome in the long term using Patient-Reported Outcome Measure Information System (PROMIS) instruments.

Patients and Methods We performed a retrospective review of all patients who underwent radial shortening osteotomy for stage 2 and 3A Kienböck's disease. Patients who had concomitant revascularization were grouped separately. We collected demographic data, data regarding type of surgery and reoperations, and radiographic data. Patient-reported outcome measures were the PROMIS Upper Extremity Computer Adaptive Testing (CAT) and Pain Interference instruments, the abbreviated Disabilities of Arm, Shoulder, and Hand (QuickDASH), and the 0 to 10 numeric rating scale for pain and satisfaction.

Results We included 48 patients who had radial shortening osteotomy alone, and 17 patients who had a combined procedure of radial shortening and direct revascularization. The rate of unplanned reoperations was 33% (16 of 48) in those who had radial shortening osteotomy and 24% (4 of 17) in those who had a combined procedure. Six (13%) of 48 patients underwent proximal row carpectomy due to failed radial shortening osteotomy. No salvage procedures were performed after combined radial shortening/revascularization. Median PROMIS Physical Function CT scores were 56 (interquartile range [IQR]: 44–56) and 56 (IQR: 41–56), respectively. Median PROMIS Pain Interference scores were 39 (IQR: 39-52) and 39 (IQR: 39-49), respectively. Median QuickDASH scores were 2.3 (IQR: 0-23) and 4.5 (IQR: 2.3-14), respectively.

Conclusion Radial shortening osteotomy for symptomatic Kienböck's disease yields reasonable long-term function. We observed that approximately one in eight patients

Keywords

- ► Kienböck's disease
- ► lunatomalacia
- radial shortening osteotomy
- vascularized bone
- patient-reported outcome
- ► long-term follow-up

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underwent salvage surgery after radial shortening, and this should be taken into account when making the initial decision to treat Kienböck's disease surgically. There appeared to be no benefit of direct revascularization in addition to radial shortening in terms of patient-reported outcome in the long term.

Level of Evidence This is a Level IV, therapeutic study.

The choice of surgery for symptomatic Kienböck's disease depends mainly on radiographic factors such as disease stage¹ and ulna variance,² as well as patient and surgeon preference.^{3,4} Radial shortening osteotomy is a widely accepted procedure for Kienböck's disease without secondary changes in carpal alignment or radiocarpal osteoarthritis (e.g., stage 2 and 3A according to the Lichtman classification) and negative or neutral ulna variance.⁵ Radial shortening osteotomy levels the distal surfaces of the radius and ulna, redistributes the axial load across the carpus, and subsequently decreases mechanical pressure on the lunate.⁶ In contrast, insertion of vascularized bone or arteriovenous pedicle grafts into the lunate aims to induce neovascularization and new bone formation in the lunate.

It is hoped that surgery can halt disease progression, decrease pain, and improve function. Some patients do not respond well enough to surgery and are offered salvage surgery, such as proximal row carpectomy or arthrodesis, due to progressive pain and carpal collapse.^{7,8} It is unclear how often patients have salvage surgery after initial radial shortening osteotomy. Furthermore, long-term patient-reported outcome data are scarce.

The aim of this study was to determine the rate of salvage surgery for failed initial treatment and any other unplanned reoperations after radial shortening osteotomy for symptomatic Kienböck's disease. Secondarily, we studied the patient-reported outcome in the long term using Patient-Reported Outcome Measure Information System (PROMIS) instruments. Finally, we studied patient-reported outcome using the more commonly used abbreviated Disabilities of Arm, Shoulder, and Hand (QuickDASH) and numeric rating scale (NRS) for pain and satisfaction.

Patients and Methods

Our Institutional Review Board approved the protocol for this retrospective study. We identified all patients who had a radial shortening osteotomy for symptomatic Kienböck's disease between January 1992 and December 2013 from our institutional database and performed medical record and radiograph review. We included patients who were at least 18 years old at the time of the study. We excluded patients who were skeletally immature at the time of surgery (n=1), had prior surgery for their Kienböck's disease (n=6), had Lichtman stage 1 (n=2), or had a radioscaphoid angle of more than 60 degrees on preoperative wrist radiographs, representing Lichtman stage 3B (n=4).

Our study population consisted of 65 patients, of whom 48 had a radial shortening osteotomy and 17 had a combined

procedure of radial shortening osteotomy and direct revascularization. The average age at the time of surgery was 37 years (interquartile range [IQR]: 29–44 years; range: 17–57 years). Thirty-seven patients (57%) were men. Hand dominance was recorded for 47/65 patients, and the dominant side was affected in 66%. In all patients, the diagnosis was made based on clinical and radiographic findings. In 13 patients, magnetic resonance imaging (MRI) scans were obtained by our institution. Some patients had outside advanced imaging. The preoperative Lichtman stages were 2 in 23 patients and 3A in 42 patients. All patients had undergone a trial of immobilization for at least 6 to 12 weeks prior to surgery.

The radial shortening osteotomy was performed using a volar approach and plating in all patients. An osteotomy was created using an oscillating saw, and, on average, 2 mm (IQR: 1-3 mm; range: 1-5 mm) of bone was resected. Internal fixation was obtained by means of a (limited contact) dynamic compression plate, T-plate, or RAYHACK system (Wright Medical Group). The median postoperative ulna variance amounted to -0.8 (IQR: -1.9 to 0.45) millimeter. Of the 17 patients (26%) who had a concomitant revascularization procedure, 13 patients had a pedicled vascularized bone graft from the distal radius to the lunate based on either the third and fourth (n=4) or fourth and fifth (n=9) extensor compartmental arteries, and 4 patients had implantation of the superficial branch of the radial artery and vein into the lunate, packed with free cancellous bone from the distal radius. Four patients had a wrist arthroscopy with synovectomy, seven patients had excision of both the anterior and posterior interosseous nerves or the posterior interosseous nerve alone, two patients had open carpal tunnel release, one patient had resection of a giant cell tumor of the thumb, and one patient had submuscular transposition of the ulnar nerve at the elbow performed in the same surgery. Postoperatively, a splint was applied for approximately 4 weeks in all patients. The procedures were performed by 11 attending orthopedic hand surgeons at two academic medical centers and one affiliated community hospital.

We reached out to all 65 patients by mail and telephone to invite them for questionnaire follow-up. Of the 65 patients, 24 did not respond or had incorrect contact information, 5 declined to participate, 4 agreed but never filled out the questionnaires, 2 did not speak English, and 2 had died of circumstances unrelated to Kienböck's disease. The 28 patients whose patient-reported outcomes were collected resembled the remaining 37, who were only available for medical record review, in terms of age, sex distribution, preoperative Lichtman stage, and additional surgeries (p > 0.05).

Study Variables and Patient-Reported Outcome Measures

The primary outcome measures were subsequent salvage procedures and any other unplanned reoperations related to the radial shortening osteotomy or Kienböck's disease. Secondary, we studied patient-reported outcomes. The main patient-reported outcome measure was the PROMIS Physical Function Upper Extremity Computer Adaptive Testing (CAT) instrument. Computerized adaptive testing optimizes the questionnaire administration by distributing only relevant items based on previous responses, shortens completion time, and decreases floor and ceiling effects. The PROMIS Upper Extremity CAT measures patient-reported disability and results in a standardized *t*-score with a mean of 50 and a standard deviation (SD) of 10. A lower *t*-score means more disability. We prefer the PROMIS instruments because of their efficiency and precision. 11

In addition, we administered the PROMIS Pain Interference instrument to measure self-reported consequences of pain on social, cognitive, emotional, physical, and recreational aspects of life. A higher score means more impediment in daily life due to pain. We also administered the QuickDASH questionnaire, which uses 11 items to measure physical function and symptoms in patients with upper extremity disorders. The QuickDASH results in a score between 0 and 100, and a higher score represents more disability. Finally, patients were asked to rate their current pain as well as their satisfaction with the surgery on a 0 to 10 NRS, and whether they would elect to have the same surgery again. Patient-reported outcome data were collected using RedCap (Research Electronic Data Capture, Vanderbilt University, Nashville, TN).

We gathered information on the following explanatory variables: age at the time of surgery, sex, affected side and hand dominance, initial Lichtman stage, date and type of surgical procedure(s), and concomitant procedures. The ulna variance in relation to the midpoint between the volar and dorsal lips of the distal radius was measured on postoperative radiographs using the method of perpendiculars. ¹²

Table 1 Patient characteristics (n = 65)

Statistical Analysis

We reported categorical data as absolute numbers and percentages, and we used the Fisher exact test to compare the proportions of categorical variables between two groups. We reported continuous data as median and interquartile range (IQR) because most data were nonnormally distributed. A Mann–Whitney *U* test was used to compare the median of continuous variables between two groups. Statistical analyses were performed using Stata 14 (StataCorp., College Station, TX), and a *p*-value of less than 0.05 was considered statistically significant.

Results

We stratified the 65 patients into those who had a radial shortening osteotomy alone (n = 48) and those who had a radial shortening combined with a revascularization procedure (n = 17; **Table 1**). Of the former 48 patients, 16 (33%) had an additional unplanned surgery (**Table 1**). In six (13%) patients, radial shortening had failed to resolve the symptoms and, subsequently, a proximal row carpectomy was performed after a median of 18 months (IQR: 7–55 months; range: 7 months to 9.3 years). Four of those patients had preoperative Lichtman stage 2, whereas two patients had stage 3A preoperatively (p = 0.17). Two (4.2%) other patients developed symptoms of ulnocarpal impaction after surgery and subsequently had an ulna shortening osteotomy. A total of seven (15%) patients had removal of the symptomatic plate and screws, and for two of those it was the only reoperation.

Of 17 patients, 4 (24%) had an additional surgery after combined radial shortening/vascularized bone grafting (\neg **Table 1**). None of these patients required salvage surgery. All four patients had removal of symptomatic hardware, of whom one patient also had arthroscopic debridement of the wrist. \neg **Table 2** provides an overview of all unplanned reoperations. The rates of reoperation were comparable for both groups (p = 0.56; \neg **Table 1**).

	Radial shortening osteotomy (n = 48)		Combined radial shortening and VBG ($n = 17$)		<i>p</i> -Value
	Median or number	IQR or %	Median or number	IQR or %	
Age, median (IQR)	35	27-42	44	31–51	0.025
Men, n (%)	31/48	65%	6/17	35%	0.048
Preoperative Lichtman					
Stage 2, n (%)	19/48	40%	4/17	24%	0.38
Stage 3A, <i>n</i> (%)	29/48	60%	13/17	76%]
Postoperative ulna variance (in millimeter), median (IQR)	-0.8	-1.8 to 0.3	-0.1	-2.6 to 0.6	0.75
Reoperation, n (%)	16/48	33%	4/17	24%	0.55
Salvage surgery, n (%)	6/48	13%	0/17	0.0%	0.33
Follow-up, n (%)	14/48	29%	11/17	65%	0.019

Abbreviations: IQR, interquartile range, VBG, vascularized bone graft. Note: Values in bold are statistically significant (p < 0.05).

Table 2 Overview of all unplanned reoperations

Type of reoperation	Radial shortening, n (%)	Radial shortening with VBG, n (%)
Proximal row carpectomy	6/48 (13%)	0/17
Hardware removal only	2/48 (4.2%) ^a	3/17 (18%) ^b
Arthroscopic debridement	3/48 (6.3%)	1/17 (5.9%)
Ulna shortening osteotomy due to new-onset ulnar-sided wrist pain	2/48 (4.2%)	0/17
Revision radial shortening due to loose hardware	1/48 (2.1%)	0/17
Repair of radius nonunion with callus autograft	1/48 (2.1%)	0/17
Repair of ruptured extensor pollicis longus tendon due to prominent hardware	1/48 (2.1%)	0/17

Abbreviation: VBG, vascularized bone grafting.

Excluding the six patients who underwent an additional salvage procedure, patient-reported outcomes were available for 25 of 59 patients, corresponding to a response rate of 42%. The median time between the primary surgery and questionnaire follow-up was 13 years (IQR: 9–14 years; range: 4–25 years). Some patients had additional procedures in the meantime (\neg **Table 2**). \neg **Table 3** summarizes and compares the average patient-reported outcomes for patients who had radial shortening alone (n=14) and those who had combined radial shortening/revascularization procedures (n=11). We observed no notable differences in any of the patient-reported outcomes between both groups. One patient indicated that she would not elect to have a radial shortening osteotomy again. There were no differences in whether the dominant or nondominant side was affected for any of the outcomes in both groups (p > 0.05).

Posthoc Power Analysis

A posthoc power analysis demonstrated that the two groups of 14 and 11 patients provided 80% statistical power (two-tailed α : 0.05) to detect a difference in average PROMIS Upper Extremity CAT score, with a large effect size of 1.18 using a parametric test. The PROMIS instruments are designed in order that 1 SD is equivalent to 10 points. In other words, we would have been able to detect a difference in PROMIS Upper Extremity CT score of 12 points or more between the groups, with 80% statistical power using a parametric test. Considering that the average minimal clinically important difference for the PROMIS Upper Extremity CAT is reported to be 9.0 in various hand conditions, 13 we would have been able to detect such a clinically relevant difference with 57% statistical power using a parametric test.

Table 3 Characteristics and patient-reported outcome of patients who were available for follow-up (n = 25)

	Radial shortening osteotomy (n = 14)		Combined radial shortening and VBG ($n = 11$)		<i>p</i> -Value
	Median or number	IQR or %	Median or number	IQR or %	
Age, median (IQR)	40	29–42	41	31–51	0.35
Men, n (%)	8/14	57%	4/11	36%	0.43
Preoperative Lichtman					
Stage 2, n (%)	4/14	29%	4/11	36%	> 0.99
Stage 3A, n (%)	10/14	71%	7/11	64%	
Reoperation, n (%)	3/14	21%	2/11	18%	> 0.99
	Median (n = 14)	IQR	Median (<i>n</i> = 11)	IQR	
Follow-up (in years)	13	9–17	13	9–14	0.69
PROMIS Upper Extremity	56	44–56	56	41–56	0.52
PROMIS Pain Intensity	39	39–52	39	39–49	0.80
QuickDASH	2.3	0-23	4.5	2.3-14	0.61
Pain (0-10)	1	0-3	1	0-2	0.40
Satisfaction (0–10)	9	7–10	10	8–10	0.36

Abbreviations: IQR, interquartile range, VBG, vascularized bone graft.

^aFive more patients had hardware removal after or during proximal row carpectomy (n = 2), arthroscopic debridement (n = 2), and nonunion repair (n = 1).

^bOne more patient eventually had hardware removal after initial arthroscopic debridement.

Discussion

This study has some shortcomings. First, although the 42% response rate is comparable to other studies, patient-reported outcomes were available for less than half of the patients. This may have introduced sampling or attrition bias. Second, this is a retrospective study of patients treated by several surgeons over a 22-year span at three hospitals. Presurgical data were not reported homogeneously in the medical record; therefore, we were unable to determine postoperative change. Third, complications and reoperations may have been underreported for patients who potentially had their follow-up elsewhere.

We observed that 6 of 48 patients had proximal row carpectomy after radial shortening osteotomy due to persistent pain. No salvage procedures were performed after combined procedures of radial shortening and vascularized bone grafting. Perhaps, additional revascularization has contributed to the lower incidence of salvage procedures, but with the numbers available we did not observe a difference. Kakinoki et al reported successful treatment of three patients with persistent pain after radial shortening osteotomy by vascularized bone grafting.¹⁴ Four patients had an additional arthroscopic debridement and no further surgeries. Previous research demonstrated that arthroscopic debridement increased wrist functional range of motion, provided excellent pain relief, and improved health-related quality of life in patients with Kienböck's disease. 15 Two of 48 patients had subsequent shortening of the ulna due to new onset of symptoms consistent with ulnocarpal impaction. Our rate does not differ from the rates reported in the literature. 8,16 Other complications were mostly related to hardware (e.g., removal of symptomatic hardware, extensor pollicis longus rupture, loose hardware).

We found that, in the long term, the average upper extremity physical function after surgery for Kienböck's disease (median: 56) was better than that of the general U.S. population (mean: 50) in terms of PROMIS Upper Extremity CAT. To place into context, patients with common hand conditions such as carpal tunnel syndrome (mean: 43), thumb carpometacarpal osteoarthritis (mean: 46), or trigger digit (mean: 47), have lower average PROMIS Physical Function scores at presentation. Truthermore, the patients in our study had low average Pain Interference scores (median: 39), mean-

ing that they had little effect of pain on activities of daily life compared with patients who presented with carpal tunnel syndrome (mean: 61), trigger digit (mean: 58), thumb carpometacarpal osteoarthritis (mean: 60), or even Dupuytren disease (mean: 52).¹⁷ These findings suggest that patients with Kienböck's disease in our study adapted well after surgery.

To compare our findings with existing literature, we also administered the QuickDASH questionnaire and NRS for pain. Normative values for the DASH and QuickDASH range from 10 to 13 in the general population. ^{18–20} The relatively low (good) average QuickDASH (median: 2.3) and pain (median: 1) scores after radial shortening are in line with those reported in the literature. Studies with more than 10 years of follow-up on average report mean (Quick)DASH scores between 6.1 and 14 and pain scores between 0.2 and 3.0 after radial shortening osteotomy (**Table 4**).^{21–25} Luegmair et al found that patients had poorer physical function when the dominant side was affected.²⁵ We found no such difference by hand dominance. Eleven patients in our study had a revascularization procedure performed in the same surgery. Patient-reported outcomes were comparable to those of patients who had radial shortening alone. Dehghani et al concluded from their randomized study that a combination of radial shortening and vascularized bone grafting is more efficient than shortening alone in terms of Mayo Wrist Score.²⁶ The follow-up in that study was only 9 months. Table 4 summarizes the literature on outcome of radial shortening osteotomy for symptomatic Kienböck's disease with average follow-up duration of more than 10 years.

Most studies report improvement of pain and good patient-reported physical function, but whether radial shortening alters radiographic disease progression is debated.²⁷ Long-term pain relief and good patient-reported outcome have also been reported after nonoperative management of Kienböck's disease. Keith et al²⁸ and Viljakka et al²⁹ report mean DASH scores of 19 and 11, respectively, in patients with Kienböck's disease after more than 10 years on average without surgery. It is unclear whether surgery contributes to the relatively good patient-reported outcomes in the literature or whether we are observing the result of the benign natural course of the pathophysiology of Kienböck's disease. Further research is needed in this area.

In conclusion, we observed that approximately one in eight patients underwent salvage surgery after radial shortening and that the overall rate of reoperation was 33%. This should be taken

Table 4 Overview of the literature

Author (year)	n	Age	Men (%)	Follow-up (range)	% Stage 3A or less	(Quick)DASH (SD)	Pain (SD)
Luegmair et al ²⁵	36	30	23 (64%)	12.1 (5.4–17.5)	100%	12 (13)	0.2 (0.6)-3 (2.3) ^a
Raven et al ²²	9	31	5 (56%)	22 (16–31)	67%	14 (23.7)	2.3 (2.9)
Rodrigues-Pinto et al ²¹	18	32	13 (72%)	10.3 (4–18)	67%	8.7 (8)	-
Viljakka et al ²³	16	32	13 (81%)	25 (20–33)	88%	6.1	0.9-3.0 ^a
Watanabe et al ²⁴	13	39	10 (77%)	21 (14–28)	77%	8.1 (8.4)	-
This study	14	40	8 (57%)	13 (4–25)	100%	2.3 (0-23) ^b	1 (0-3) ^b

Abbreviations: DASH, Disabilities of Arm, Shoulder, and Hand; SD, standard deviation.

^aPain during rest and during activity.

^bMedian and interquartile range.

into account when making the initial decision to treat Kienböck's disease surgically. The long-term patient-reported outcomes of radial shortening—whether or not combined with a vascularized bone graft to the lunate—for symptomatic Kienböck's disease appear to be reasonable. It is unclear whether these surgeries alter the natural history of Kienböck's disease; however, in general, these surgeries appear to yield reasonable long-term function. There appeared to be no benefit of direct revascularization in addition to radial shortening osteotomy in terms of patient-reported outcome.

Ethical Approval

This study was approved by Partners Human Research Committee, Boston, MA.

Note

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Conflict of Interest None declared.

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