



# Periprosthetic humerus fractures: classification, management, and review of the literature

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**Abstract:** Complications of shoulder arthroplasty can present significant treatment challenges and complexity especially in the setting of periprosthetic humeral fractures. This paper will provide a review of timing and location of these fractures, classification, imaging modalities to best assess both the implant stability as well as fracture pattern, treatment strategies, and results. The goal in the management of this complication is to restore component stability and maximize shoulder arthroplasty implant survivorship as well as patient functional outcomes.

**Keywords:** Shoulder arthroplasty; complications; periprosthetic fracture; revision shoulder arthroplasty

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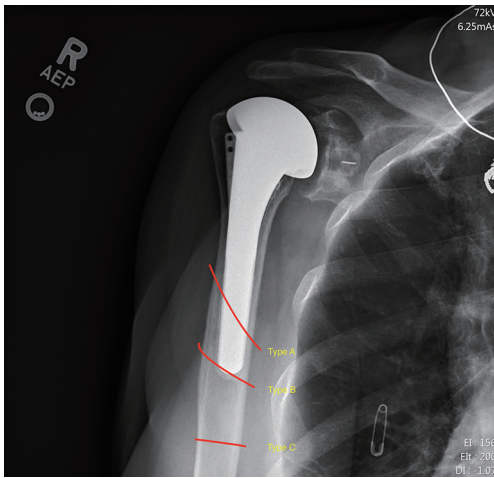
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## Introduction

Total shoulder arthroplasty (TSA) has been accepted as a reliable and predictable treatment option in the management of end-stage glenohumeral joint disease, with the frequency of the procedure increasing 2.5 folds over the past decade (1-3). Current evidence shows that primary, anatomic TSA improves both pain and function in up to 90–95% of patients (4-6). More recently, reverse total shoulder arthroplasty (rTSA) has also been shown to be effective in treating rotator cuff arthropathy, as well as failure of primary anatomic TSA (7-15). Long-term data demonstrate that these improvements in pain and function persist, and survivorship of these implants at ten years and twenty years is over 90% and 80%, respectively (10,16,17).

Despite this promising data, a wide-array of complications has been reported in the literature. Total complication rate for TSA/rTSA ranges from 3.6% to 20%, with one study examining rTSA noting a 68% complication rate (6,18-22). These complications include, but are not limited to, infection, instability, periprosthetic humerus

fractures, neurovascular injury, component loosening, rotator cuff tear, and deltoid injury (20,23,24). Anatomic TSA complications include glenoid component loosening, osteolysis around either the humeral or glenoid component, instability from subscapularis failure all of which can result in significant bone loss. Additional complications seen more often in the setting of rTSA include baseplate failure, scapular/acromial stress fractures, instability, and component dissociation (14,15,25). Scapular notching is often frequently reported as a complication following rTSA, but the clinical significance of this is still unknown as there has been no study to demonstrate correlation with severity of notching and patient reported outcomes. While earlier data suggested a higher overall complication rate in rTSA when compared to anatomic TSA, more recent evidence suggests this may not be the case (13,26). The majority of investigations reporting complication rates following rTSA included a heterogeneous cohort of patients with a variety of preoperative shoulder pathology including massive rotator cuff tears/rotator cuff tear arthropathy, primary osteoarthritis, post traumatic arthritis, revision shoulder



**Figure 1** Wright and Cofield classification system (43).

arthroplasty, inflammatory arthritis, and fracture (27). Additionally, Jacxsens *et al.* (28) demonstrated that the significant amount of heterogeneity with regards to classifying and reporting complications makes comparison difficult, and a standardized scheme would be beneficial. The incidence of complications requiring reoperation is 5–11% and has been shown to be more frequent following revision procedures (18,29–31). A recent review article by Bohsali *et al.* reports an overall downward trend in complications over the last decade overall (27).

Periprosthetic humerus fractures are of particular importance, as they frequently lead to the need for revision surgery (32,33). These fractures can take place intraoperatively or postoperatively, occurring 0.9–3.5% and 1.0–3.0% of the time, respectively (20,21,25,27,29,32,34–38). They are more frequent during revision procedures, especially during extraction of a well fixed humeral stem (27,39–42). Preparing the humerus for an implant renders the bone more susceptible to fracture, and evidence shows that periprosthetic humerus fractures have a higher nonunion rate than native humerus fractures (6,43,44). This review, focuses on classification systems, risk factors, non-operative as well as operative management strategies, complications, and outcomes for both intraoperative and postoperative periprosthetic humerus fractures in the setting of anatomic and rTSA.

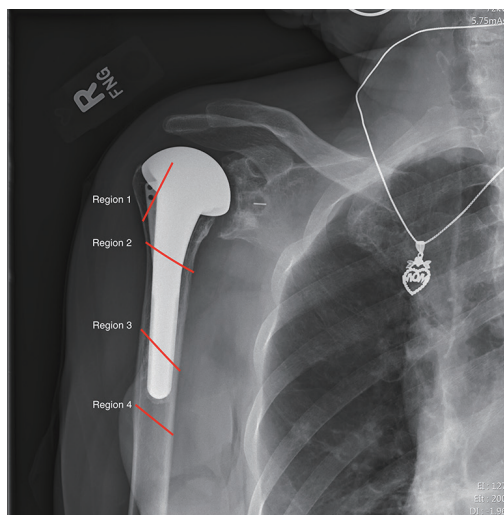
## Classification

Wright and Cofield were the first to devise a classification system for postoperative periprosthetic humerus fractures

in 1995 (43). They reported on a series of nine patients, and each was classified based on fracture pattern in relation to the distal tip of the prosthesis (*Figure 1*). While both types A and B fractures were centered at the tip, they differed in the amount of proximal extension. Type C fractures involved the humeral shaft distal to the tip of the implant. Type A fractures required surgery, with the fixation strategy dependent on implant stability. Long oblique or spiral type B fractures could be managed without surgery, while transverse or short oblique fractures fared better with operative intervention. Finally, type C fractures could be managed similarly to shaft fractures in the native humerus. This scheme has been shown to have good intraobserver (mean kappa, 0.69; range, 0.52–0.89) but poor interobserver reliability (mean kappa, 0.37; range, 0.24–0.50) (45). In 1999, Worland *et al.* published a similar classification scheme based on six patients, but subdivided type B fractures into B1–B3 based on fracture pattern and implant stability (46). B1 fractures are spiral fractures with a stable prosthesis, B2 are short oblique or transverse fractures around the tip of the stem and are also stable, and B3 are fractures resulting in an unstable stem. They noted that types A and C behave similar to native humerus fractures, while type B fractures present more of a treatment challenge depending on the stability of the stem. Groh *et al.* presented a similar classification system in 2008 developed from 15 patients which was also entirely based on the location of the fracture (47). Type I fractures occurred at the tip of the prosthesis, type II the fracture extended from proximal humeral shaft to beyond the distal aspect of the stem, and type III were fractures distal to the tip of the prosthesis.

Campbell *et al.* classified fractures as belonging to one of four regions (*Figure 2*) (36). They assumed that fractures in the proximal humeral metaphysis have different implication for both healing and prosthetic stability than those occurring in the diaphysis. Of the 21 fractures in their series, 16 occurred intraoperatively. They concluded that intraoperative fractures all required anatomic reduction and stable intramedullary fixation. Region 1 and 2 fractures can be treated with standard length prosthetic stem and supplemental suture or cerclage wire fixation. Region 2 and 4 fractures required conversion to a long-stem implant with supplemental cerclage wiring.

Duncan *et al.* presented a classification system that could be broadly applied to any periprosthetic fracture which they entitled the “Unified Classification System” (48). This classification is based on whether the implant is well fixed, the patient’s bone quality, location with relation to soft



**Figure 2** Campbell classification system (36).

tissue attachments, and whether the bone supports two joint replacements (i.e., a periprosthetic fracture between a total elbow and TSA). Its clinical utility in the classification and management of periprosthetic humerus fractures has not been well-elucidated in the literature as of this time.

The most comprehensive classification system for periprosthetic humerus fractures to date was published in 2016 and validated in 2018 (49,50). Kirchoff *et al.* aimed to create a structured approach to classifying periprosthetic humerus fractures that could be used to guide treatment in an algorithmic fashion (Figure 3A,B). Unlike previous iterations, this system takes into account the type of humeral prosthesis (stemless *vs.* anatomic *vs.* reversed), status of the rotator cuff, location of fracture, fracture pattern, and implant stability.

## Risk factors

### Patient factors

There have been several identified risk factors for both intraoperative and postoperative periprosthetic humerus fractures during TSA/rTSA. Intraoperative fractures most commonly occur during stem extraction during revision and most often involve the greater tuberosity, as this bone is often thin from stress shielding in the presence of a prior implant. Glenoid exposure in a shoulder that is severely retroverted or severe soft tissue contracture places the humeral shaft at increased risk for spiral diaphyseal fracture. Finally, overstuffing the humeral canal with a

press fit humeral stem can result in metaphyseal fractures of the proximal humerus during impaction of the implant. Singh *et al.* performed a retrospective case series identifying female sex and posttraumatic glenohumeral arthritis as two risk factors for intraoperative periprosthetic humerus fractures (40). The authors hypothesized that the higher prevalence of osteoporosis in elderly females could explain this finding. Prior hemiarthroplasty and a history of instability have also been shown to increase risk of intraoperative humerus fractures (51). Higher number of medical comorbidities, increasing age, female sex, osteopenia, avascular necrosis, and rheumatoid arthritis are other risk factors that have been associated with an increased likelihood of postoperative humerus fractures (6,25,32,34,36,43,52-56).

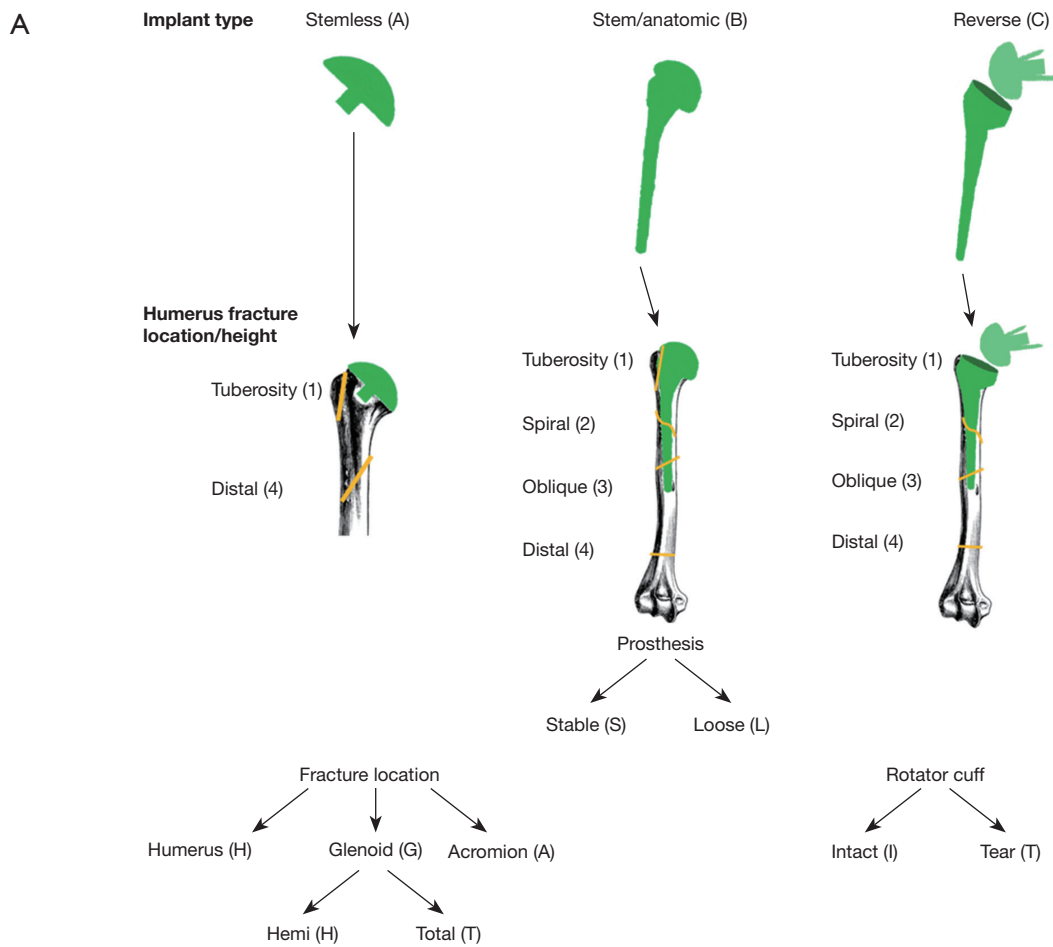
### Intraoperative factors

In addition to patient related factors, which often cannot be modified, there are multiple technical considerations a surgeon must be aware of in order to decrease the risks of a periprosthetic humerus fracture. Initially, it was believed that cementless implants conferred a high probability of fracturing the humerus, similar to press-fit femoral implants, but most recent evidence reveals no difference between cemented and cementless prostheses (37,40,57,58). Avoiding: (I) excessive external rotation of the humerus during reaming and/or broaching; (II) overzealous reaming (ream by hand); (III) under-reaming; (IV) malalignment of the stem; and (V) inserting oversized broach/prostheses are important technical aspects which have been shown to increase the risk for fracture (6,29,36,54,58). Appropriate patient positioning, ensuring adequate exposure with adequate soft tissue and capsular releases are imperative in avoiding these fractures, with some evidence hinting that an anterosuperior lateral approach is safer than the more traditional deltopectoral approach when performing rTSA—although this remains controversial (6,34,59). In revision procedures, periprosthetic humerus fracture frequently occurs either during implant removal or insertion; the remainder being during reaming or broaching (51,60). Most commonly the greater tuberosity is fractured during stem extraction or exposure of the glenoid.

## Clinical evaluation and management

### Intraoperative fractures

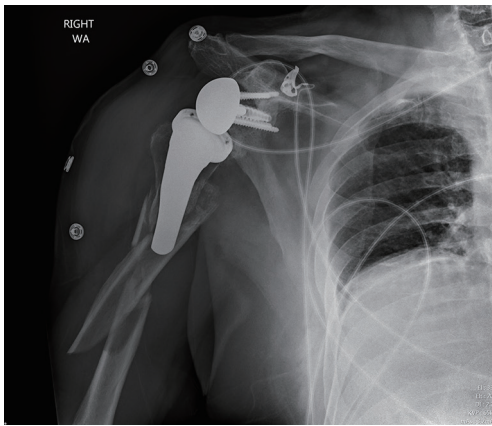
We will begin by reviewing the management strategies



**B**

Type	Treatment	Type	Treatment	Type	Treatment
A-A-X-X-X	Conservative vs ORIF	B-A-X-X-X	Conservative vs ORIF	C-A-X-X	Conservative vs ORIF
A-G-H-I-X-X	Conservative vs ORIF vs TEP	B-G-H-I-X-X	Conservative vs ORIF vs TEP	C-G-X-L	Revision glenoid
A-G-H-T-X-X	Convert to reversed	B-G-H-T-X-X	Convert to reversed	C-H-1-S	Conservative vs ORIF
A-G-T-I-X-S	Conservative	B-G-T-I-X-S	Conservative	C-H-2-S	ORIF
A-G-T-I-X-L	Revise glenoid	B-G-T-I-X-L	Revision glenoid	C-H-3-S	ORIF
A-G-T-T-X-X	Convert to reversed	B-G-T-T-X-X	Convert to reversed	C-H-X-L	Convert to reversed long stem +/- ORIF
A-H-H-I-1-S	ORIF	B-H-H-I-X-S	ORIF		
A-H-H-I-1-L	Convert to anatomic stem	B-H-H-I-X-L	Revise to TEP + new stem		
A-H-H-T-1-L	Convert to reversed	B-H-H-T-X-S	Convert to reversed		
A-H-H-T-4-L	Convert to reversed + ORIF vs reversed long stem	B-H-H-T-1-L	Convert to reversed		
		B-H-H-T-2-L	Convert to reversed + ORIF		
		B-H-H-T-3-L	Convert to reversed + long stem		
		B-H-H-T-4-L	Convert to reversed long stem + ORIF		

**Figure 3** Algorithm representation of Kirchhoff classification system as well as description of how the classification system can direct treatment. (A) Kirchhoff classification system (49,50); (B) treatment recommendations based on classification.



**Figure 4** Radiograph of right shoulder periprosthetic humerus fracture.

for intraoperative periprosthetic humerus fractures. We elected to discuss this separately as, assuming the fracture is identified during surgery, its treatment algorithm often differs significantly when compared to fractures discovered during follow up. Once a periprosthetic humerus fracture has occurred during surgery, several factors must be considered to determine the best strategy on how to proceed. These factors include location of fracture, fracture displacement, and bone quality. The goal is to employ techniques of rigid internal fixation, to prevent the fracture affecting the patient's postoperative rehabilitation (58). Postoperative changes in rehabilitation are most significantly influence by fracture location, those involving the greater tuberosity and rotator cuff insertion need to proceed cautiously to allow for healing without displacement by the pull of the tendons. Fractures more distal involving the metaphysis or diaphysis may allow for immediate mobilization as would occur in a primary setting as often times a stable, rigid construct with long humeral stems supplemented with cables or suture can be achieved.

For nondisplaced fractures of the greater tuberosity that do not extend distally, observation alone is often adequate (6,37). If there is any displacement, internal fixation with suture or cerclage wiring is recommended (25,47). Normal postoperative rehab protocol can then be utilized for these patients. Surgical fixation is required if the tuberosity fracture extends into the proximal aspect of the humeral shaft or if it involves the surgical neck. If the bone quality is deemed reasonable, treatment with either suture or wire fixation and revision to long stemmed component or standard length prosthesis (if it extends at

least 3 cortical diameters past fracture site) combined with inter-fragmentary screws is adequate (36). Bone graft can be utilized in cases of bone loss (37).

Treatment of shaft fractures depends on its location relative to the tip of the prosthesis. For fractures proximal to the tip of the prosthesis, revision to long stem implant, which extends at least two cortical diameters past the fracture site, with cerclage wire or cable fixation with or without allograft struts is recommended (6,36,37,47,58). Fractures distal to the prosthesis, but the prosthesis is deemed unstable, the same management strategy applies. If the prosthesis is deemed stable intraoperatively, the implant can be retained and internal fixation with hybrid plate/cerclage wire construct can be utilized (47,58).

### *Postoperative fractures*

#### **Preoperative evaluation**

A careful history and physical examination should be performed on every patient presenting with a periprosthetic humerus fracture. A majority of these are caused by low-energy mechanisms such as fall from standing and determining pre-injury functional status can help guide subsequent treatment. It is important to determine patients whether or not pain was present prior to the fracture, as that could be a sign of implant loosening or low virulence infection such as propionibacterium acnes. Although inflammatory markers such as ESR/CRP are valuable in evaluating for an occult infection, these studies are often unreliable in the setting of an acute fracture. Presence of bacteria can be better determined at time of revision surgery, but significance of unexpected positive cultures continues to be investigated. Medical records are helpful in determining preoperative range of motion, identifying surgical approach, as well as confirming which implants are present. Knowledge of what implant is being revised is helpful for preoperative planning as well as nuances of the system with regards to extraction. Physical examination is often pain-limited, but the condition of the skin, prior scars, as well as neurovascular status should be carefully evaluated and accurately documented. It is important to evaluate the axillary and radial nerves specifically as these neurologic structures are at particular risk for injury depending on the location of the periprosthetic fracture and zone of injury.

AP, true AP, scapular-Y, and axillary lateral views (*Figure 4*) of the shoulder should be obtained, in addition to full length AP and lateral humerus XRs (*Figure 5*). CT scan is indicated to better determine extent of the fracture

as well as evaluating humeral and glenoid bone stock (56). It also provides valuable information regarding glenoid morphology and remaining glenoid vault. CT arthrogram or ultrasound can be helpful to assess the status of the rotator cuff and CT arthrogram can provide additional details on stability of the components especially the glenoid in the setting of anatomic TSA.

### Nonoperative management

Nonoperative management has been shown to be successful when appropriately indicated (*Figure 6A,B*). Multiple studies show that fractures distal to a well-fixed prosthesis (type C) can be treated nonoperatively similar to native humeral



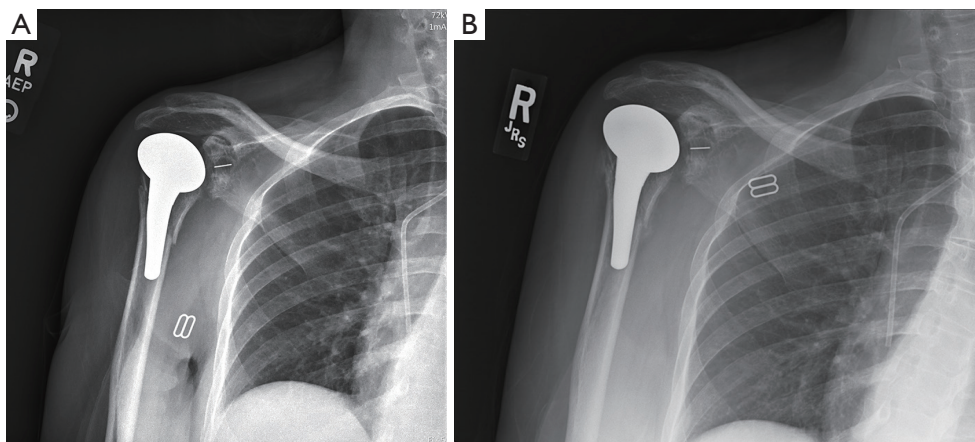
**Figure 5** Full length humerus radiograph right periprosthetic humerus fracture.

shaft fractures (43,61-64). Additionally, if the fracture is proximal to the tip of a well-fixed implant, a trial of non-operative management is reasonable—although this remains controversial (33,43,61). Acceptable closed reduction parameters are similar to native humerus fractures (63). However, if the prosthesis is loose, surgical intervention is indicated—assuming the patient is medically fit for surgery.

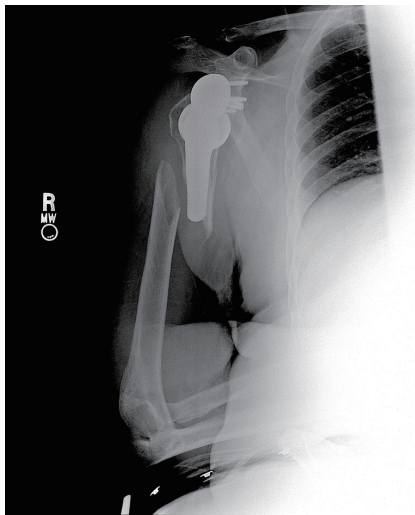
### Operative management

While heterogeneity remains in the literature, commonly agreed upon indications for surgical treatment of periprosthetic humerus fractures are displaced/unstable fractures and fractures around a loose humeral component (45,49,65).

The first step when planning treatment is to determine implant stability. Preoperative XRs can be evaluated using criteria presented by Sperling *et al.* and Sanchez-Sotela *et al.* (66,67). They determined that humeral implants with >2 mm of lucency in at least 3 of 8 humeral zones or those where 2 of 3 independent observers believed there to be subsidence or tilt of the implant were at increased risk for loosening (*Figure 7*). However, the most accurate method to determine implant stability is intraoperatively. This can be performed by applying a direct force to the humeral stem tip after exposure of fracture site (45,68). If equivocal, proximal humerus exposure can be performed by taking down subscapularis tendon and capsule, and the bone-implant interface can be directly observed while again applying pressure to the stem tip through the fracture site. Alternatively, the rotator interval can be opened proximally to facilitate the evaluation of the stem at the level of the



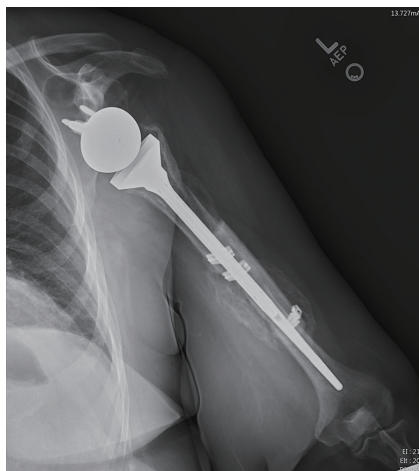
**Figure 6** Radiographic example of a stable periprosthetic fracture treated conservatively. (A) Right shoulder radiograph stable periprosthetic fracture; (B) right shoulder radiograph demonstrating well healed fracture following conservative management.



**Figure 7** Radiographs right shoulder periprosthetic fracture with loose unstable humeral prosthesis.



**Figure 9** Right shoulder radiographs of periprosthetic fracture treated with long stem humeral prosthesis supplemented with allograft struts and nylon cables.



**Figure 8** Radiographs of left humerus revised to long stem implant supplemented with nylon cables and strut allograft which is well healed and incorporated.

glenohumeral joint (68).

Recent data show that type A fractures can be treated nonoperatively if they are nondisplaced (25,29). However, many believe these fractures should be managed with revision to long stem prosthesis with additional fixation determined by bone quality (plate, cerclage wiring) (63). Successful operative management of these fractures with just suture fixation without implant revision has been reported (20,29). Type B fractures with a stable implant can be treated with implant retention and osteosynthesis

(25,33,65,68-70). Specific constructs vary widely in the literature, without significant difference in union rates. If the implant is loose intraoperatively, conversion to long stem prosthesis is treatment of choice. Implant should span 2 or 3 cortical diameters past the fracture site (33). Supplemental fixation with cerclage wire with or without allograft strut augmentation or plate constructs is often helpful (Figure 8). Augmentation with allograft is also an option (45). When there is significant proximal humeral bone loss from either the fracture, stem extraction or both proximal humerus composite allograft can be used. Type C fractures with a loose stem should be treated similarly. Type C fractures with a well-fixed stem can be treated similarly to native humeral shaft fractures. Of note, surgeons will likely have to utilize cables, wires, unicortical screws, if locked plating is used in these settings due to the presence of the prosthesis proximally or the humeral stem can be revised to a long stem construct (47,49,64,70-76) (Figure 9).

## Outcomes

Overall union rate after surgery for periprosthetic humerus fractures is high, with one series reporting 97% union rate in 36 patients treated either with revision arthroplasty or ORIF (45). In the ORIF group (n=17), fractures healed at an average of 6.8 months. Of the 6 that had pre-fracture ASES scores, 5 returned to that level postoperatively. In the

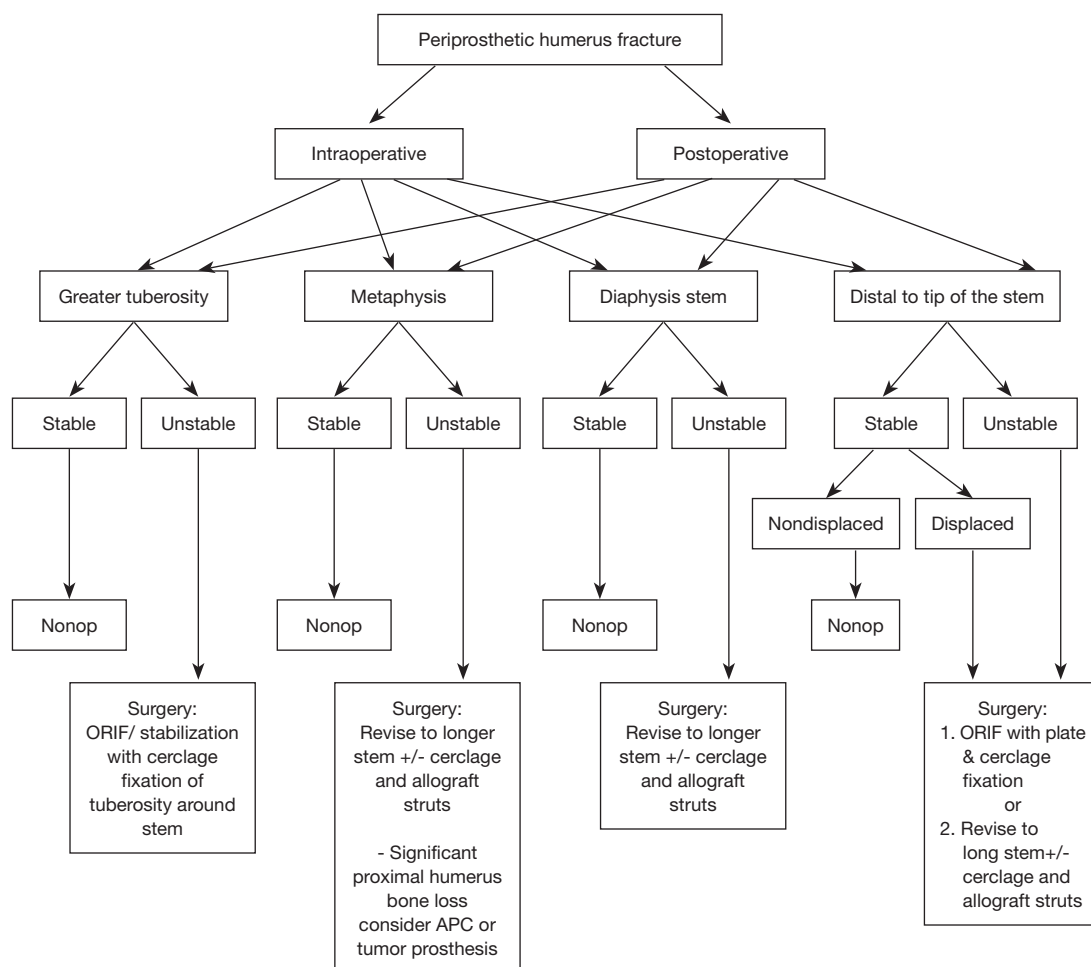


Figure 10 Treatment algorithm for periprosthetic humerus fractures.

revision arthroplasty group (n=19), average time to union was 7.7 months. There was a statistically significant increase in preoperative versus postoperative ASES scores. Despite this promising date, they noted a 39% complication rate and 19% reoperation rate. Other studies similarly show high complication rates and note that patients do not do nearly as well as they did after their index procedure (43,53,56).

**Summary/authors recommendation**

Timing, location, and implant stability of periprosthetic humeral fractures related to shoulder arthroplasty dictates treatment recommendations. While the published outcomes are limited to retrospective case series for evidence-based treatment decisions, we have developed a simple treatment algorithm to allow for best predictable functional outcomes (Figure 10).

Intraoperative humeral fractures occurring either primary or revision shoulder arthroplasty most commonly involve the greater tuberosity or humeral metaphysis. (I) Fractures of the greater tuberosity are addressed with high strength non-absorbable doubled wracking hitch suture technique where the suture is passed at the tendon bone junction of the supraspinatus/infraspinatus and secured around the humeral stem. (II) Metaphyseal humeral fractures typically occur during stem insertion or humeral preparation and can be treated with cerclage technique around the metaphysis.

Postoperative periprosthetic fractures can occur at any location along the humerus and as all of the classifications have outlined the most important factors include location and implant stability. We have simplified this into categories: can the stem be preserved, or does it need to be revised. A CT scan is always obtained and allows for perioperative assessment of remaining humeral bone as



well as fracture pattern. (I) Proximal fractures with a stable implant are treated nonoperatively. (II) Fractures involving the metaphysis/diaphysis region with in the area of the stem are driven by remaining proximal humeral bone, stem stability, and method of fixation of primary stem (i.e., how destructive would it be to the remaining bone to extract and revise the prosthesis). (III) Stems that are grossly loose are revised to long stem humeral prosthesis with cerclage supplementation around the fracture with or without allograft struts. (IV) Stems that are stable are driven by fracture pattern, and how much of the stem is involved with the fracture.

Additional outcomes-based studies are needed to provide data on how these fractures effect implant survivorship and patient reported functional outcome measures.

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*Conflicts of Interest:* MK is a consultant and receives research support from Tornier/Wright Medical both of which are unrelated to the content of this work. The other authors have no conflicts of interest to declare.

*Ethical Statement:* The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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