The vertical humeral osteotomy for stem removal in revision shoulder arthroplasty: results and technique

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Hypothesis: Revision shoulder arthroplasty represents a complex and difficult problem for the treating surgeon, with multiple potential complications. In the setting of a well-fixed humeral component, removal can lead to fractures and compromise the outcome of the revision. The current study describes and evaluates the results of a novel vertical humeral osteotomy (VHO) for stem extraction. We hypothesized that the VHO will enable successful stem extraction without perioperative or postoperative fractures.

Materials and methods: Twenty-seven patients were retrospectively identified who had a VHO for revision shoulder arthroplasty, with 23 patients available for final follow-up. Records and radiographs were reviewed for postoperative complications. Final follow-up was completed with the inclusion of shoulder scores.

Results: There were no perioperative or postoperative fractures on clinical examination and radiographic review at an average follow-up of 41 months. Average American Shoulder and Elbow Surgeons (ASES) score was 64.7 (contralateral ASES, 76.9), average Simple Shoulder Test was 6.3, and the visual analog score pain average was 1.3. There were no instability events.

Discussion: The glenoid is the more common site for failure in both hemiarthroplasty and total shoulder arthroplasty. This can lead to a difficult revision procedure if the ingrown or cemented humeral stem requires removal.

Conclusion: In the current study, we found the VHO was an effective tool for the removal of the humeral prosthesis with no perioperative or postoperative fractures.

Level of evidence: Level IV, Case Series, Treatment Study.

Keywords: Shoulder; arthroplasty; revision; humeral; prosthesis

Shoulder arthroplasty has experienced an exponential increase in available implants, indications, and techniques in contemporary orthopedics. Good to excellent results have been reported with both hemiarthroplasties and total shoulder replacements for the treatment of proximal humerus fractures, end-stage degenerative arthritis, and rotator cuff arthropathy. However, as the frequency of primary shoulder arthroplasty increases, the number of failures and required revisions will also grow.

Revision shoulder arthroplasty represents a complex and difficult problem for the treating surgeon. The extent to which component removal is necessary depends on the mode of failure. Failures can manifest due to glenoid erosion, glenoid component loosening, instability, infection, component malposition, and rarely, humeral component...
loosening. In the context of revision surgery with a well fixed humeral component that requires removal, the procedure can be extremely challenging, with significant complications. Extensive bone ingrowth and a large, intact cement mantle contribute to the difficulty with extraction of the humeral stem. However, the thin cortical bone of the humerus makes it difficult to create a safe window or L-shaped osteotomy. Resultant fracture or loss of tuberosity integrity can lead to severe postoperative dysfunction. Thus, without a safe and reliable technique for stem removal, the proximal humerus may be unnecessarily fractured or denuded of bone stock.

Sperling et al has previously described an anterior or medial cortical windowing technique to facilitate humeral stem removal. However, they reported a 20% rate of intraoperative fracture associated with this procedure and noted that, with refinement, further techniques could be developed that significantly lowered this rate. Subsequently, we have developed a vertical humeral osteotomy (VHO) technique to remove both cemented and uncemented humeral components, as described previously. The technique was designed to allow component removal without significant damage to the proximal humerus and to avoid distal windows, thus allowing reimplantation without the need for a longstem implant.

The purposes of the current study were to describe the VHO technique, report the perioperative complications, and evaluate the longer-term follow-up results. We hypothesized that the VHO would enable successful stem extraction without perioperative or postoperative fractures.

Materials and methods

We retrospectively identified 27 patients (16 women, 11 men) who required a VHO for humeral stem removal in a revision arthroplasty. Of these, 23 were available for final follow-up: 2 had died of unrelated causes, and 2 could not be located. Patients were clinically evaluated at postoperative intervals of 1 week, 1, 3, and 6 months, 1 year, and then at yearly intervals. Postoperative radiographs were obtained in the true anteroposterior (AP), lateral, and axillary planes at 3 months, 6 months, 1 year, and then annually thereafter. Clinical scores were obtained and recorded at final follow-up, including the American Shoulder and Elbow Surgeons (ASES) score, Simple Shoulder Test (SST), and visual analog scale (VAS) for pain. Radiographs from each follow-up visit were reviewed for fractures or component loosening. Finally, all operative reports were analyzed for notations of fracture, complication, or loss of tuberosity integrity.

The average follow-up was 40.9 months (range, 24-98 months). The average age at revision was 69 years (range, 52-81 years). There were 14 cemented stems and 9 uncemented humeral stems. Seven arthroplasties were initially infected, but the humeral stem (5 cemented, 2 uncemented) was not loose, therefore requiring a VHO for removal.

The presenting etiology for the original arthroplasty was fracture in 10 patients, cuff tear arthropathy in 7, and osteoarthritis in 6. The subsequent revision arthroplasties included hemiarthroplasty to reverse in 14 patients, hemiarthroplasty to total shoulder arthroplasty in 6, total shoulder arthroplasty to reverse in 1, and hemiarthroplasty to hemiarthroplasty in 2. In the 7 infections, a spacer was placed for 6 to 8 weeks, and the secondary exchange was performed by removing the spacer and implanting the new component. The VHO had been performed to remove the primary stem and was not disturbed at the secondary reimplantation.

Surgical technique

The VHO procedure is performed under scalene regional and general anesthesia in the beach chair position. The previous deltopectoral incision is used in developing the deltopectoral interval; occasionally, this will need to be extended. Dense scar tissue should be released from the undersurface of the deltoid and proximal humerus. If the subscapularis tendon is still present, it is released off the lesser tuberosity and reflected medially. To enhance humeral exposure, abundant scar tissue and remnant glenohumeral capsule are released from the anterior, inferior, and posterior glenoid rim. The axillary nerve is palpated and protected during this step. After this release, the proximal humerus is easily delivered into the open surgical wound with flexion and external rotation of the arm.

The VHO is then used to remove a well-fixed humeral component, either cemented or uncemented. This allows the surgeon to “debond” the humeral stem from the cement mantle without having to go distal to the stem tip. A small osteotome is first used around the top of the prosthesis to interrupt the interface between the implant and the tuberosity bone (Fig. 1). This is an important step to eliminate adhesion of the thin tuberosity bone to the implant or cement mantle. The prosthetic head is removed, and a 0.25-inch curved osteotome is tapped down around the proximal aspect of the humeral component circumferentially.

Cautery is used to expose the humerus vertically, beginning just lateral to the biceps groove and extending distally between the anterior deltoid and lateral pectoralis tendon insertions. This extends approximately 10 cm distally on the humerus (Figs. 2 and 3). A MicroAire oscillating saw (MicroAire Series 1000, Micro-Aire, Charlottesville, VA) is used to create a linear unicortical osteotomy along this vertical line, perforating both the cortex and underlying cement mantle down to the implant. This type of saw has a small blade that is easily controllable and makes a thin cut into the bone. The osteotomy is extended distally to just below the deltoid insertion but not below the tip of the implant.

Next, a series of osteotomes are used to gently “flex” open the humeral shaft at the osteotomy. Care is taken to avoid fracturing the opposite cortex (Fig. 4). The osteotomes are placed vertically within the osteotomy (perpendicular to the shaft) and gently twisted to open the humeral envelope. Gently repeating this “open booking” of the unicortical osteotomy on several occasions creates a visible gap between the cement mantle and prosthesis. There is no need to create an L-shaped cortical flap or window.

Once this gap is visualized, a footed impactor is placed on the medial neck of the proximal aspect of the humeral implant (Fig. 5). A mallet is used on the footed impactor to extract the humeral stem. Once the implant starts to move, care is taken to make sure the tuberosity bone is not fixed to the implant. We have found that implant-specific slap-hammers are less reliable than this technique.
After the implant is removed, a portion of the remaining cement mantle may be loosened from the surrounding cortex. Additional gentle “open booking” of the cortex facilitates removal of this mantle with osteotomes and a rongeur to a stable remnant mantle. If the need for revision is not an infectious process, then the entire cement mantle does not need to be removed. A new implant can be cemented into the old cement mantle. In the setting of an infection, all foreign material is removed, including the entire cement mantle and the distal plug. This can be done without extending the osteotomy through the use of an ultrasonic cement remover (Ultra-Drive, Biomet, Warsaw, IN) and with small bone hooks for extraction of the plug. Throughout the extraction process, the humeral shaft remains intact, without propagation of the osteotomy.

Two looped 18-gauge Luque wires are then passed circumferentially around the humeral shaft using a wire passer and spaced evenly across the proximal-distal expanse of the osteotomy (Fig. 6). A cable twister is used to lightly tighten each cerclage construct, with an assistant digitally palpating the inner surface of the osteotomy to maintain an anatomic diaphyseal reduction. A metaphyseal reamer is then used to prepare the canal by hand, with most of the cement mantle left in place. A standard length, canal, and mantle filling trial stem can then be inserted to protect the humeral shaft during glenoid preparation and component insertion (Fig. 7).

**Results**

Twenty-three patients were available for follow-up at a mean of 41 months. All revision stems were cemented: 9 were cemented into existing cement mantles, and the other 14 were cemented into the diaphyseal tube. Seven of these were a primary exchange of an uncemented stem to a revision cemented standard length stem. The other 7 were secondary re-implantations after spacer removal for infection treated by implant removal by VHO. All revision stems were standard length, with no long stem components being necessary.

No distal window or L-shaped diaphyseal osteotomies were required to remove the primary stems. There were no intraoperative diaphyseal or metaphyseal fractures. Follow-up radiographs revealed no implant-cement lucencies or loosening at an average of 41 months and no extruded cement was seen in the diaphyseal region. No periprosthetic fractures have occurred. There were no cerclage wire complications. No implant specific extractor devices or slap hammers were used with the VHO technique.
Shoulder function outcome scores were variable. At an average follow-up of 41 months, average ASES score was 64.7 (contralateral ASES, 76.9), average SST was 6.3, and the VAS pain average was 1.3. There were no instability events. All patients stated they would undergo the procedure again and were pleased with the result.

Discussion

The continued growth and refinement of shoulder arthroplasty will also result in an increase in revision surgery. Multiple authors have shown that excellent results can be attained; however, the techniques required can be complex. The etiology of failure is an important component when planning the revision procedure. The glenoid is the most common site for malfunction through loosening of the component or progression of arthrosis on the native articular surface. Less commonly, the humerus can be involved with component malposition, instability, infection, fracture, and, atypically, humeral component loosening. Thus, a well-fixed humeral prosthesis creates a difficult problem and emphasizes the importance of a safe and reliable means of stem removal. This fact was highlighted by Wall and Walch in their review of reverse shoulder replacements, “during thirteen (24.1%) of the fifty-four revision procedures, a humeral fracture occurred during removal of the primary prosthesis or cement mantle.”

The concept of an osteotomy for removal of a prosthesis has been well established in the revision total hip literature in the form of an extended trochanteric osteotomy. This has proven to be a successful tool in removing cemented and uncemented femoral components during revision procedures. However, an analogous technique for revision shoulder arthroplasty has received little attention:

- Carroll et al and Peterson et al alluded to an osteotomy procedure for extraction of a humeral stem, but neither described the technique.
- Sperling et al reported using an anterior or medial cortical window to access the humeral component. The window was resected then replaced, secured, and supplemented with allograft in 13 of 16 patients. Of note, they reported a 20% intraoperative fracture rate.
- Gohlke and Rolf described a vascularized humeral window technique based on the pectoralis insertion. They reported no loosening at final follow-up in 34 patients, with 8 complications including dislocation, infection, and a fracture in the postoperative period. However, this technique requires 2 osteotomies with a technically difficult medial osteotomy under the insertion of the pectoralis insertion. This potentially creates 2 issues: complications with the replacement and healing of the “window” piece as well as a difficulty in successful reproduction of the medial osteotomy.

Overall, the shortcomings of the reported techniques led to the development and refinement of the VHO technique.
In the current study, the VHO represents a unique osteotomy procedure for revision shoulder arthroplasty that has demonstrated no iatrogenic fractures in this case series. Twenty-three patients had good outcomes at an average of 41 months, with no perioperative or postoperative fractures. The wide range of outcomes scores is the result of a variety of failure etiologies in patients with a spectrum of functional demands.

However, there are significant limitations to the current study. The functional demands of the patients may not be representative of every patient population. A patient with high functional demands may have a higher propensity for fracture. Also, these patients were identified retrospectively and complications were not collected prospectively. Although the operative reports and clinic notes were thorough, the potential for undocumented complications does exist. However, the senior author (G.N.) has used this technique for more than 10 years, and these patients are contained within a solitary practice.

Theoretically, the described humeral osteotomy works by releasing hoop stresses in the proximal humeral shaft. This greatly loosens the stem and facilitates removal of a well-fixed component without propagation or iatrogenic fracture elsewhere in the humerus. We have used this technique with press-fit and proximally coated trabecular metal stems, as well as with cemented stems, and enjoyed a similar ease of removal. The supplemental cerclage wire fixation of the osteotomy provided adequate stability to the cortex and did not detrimentally influence the radiographic and clinical outcomes. We also have not found any problem with the extended soft-tissue exposure necessary for the osteotomy. It is easily performed through a standard deltopectoral approach and preserves the pectoralis major insertion on the medial side and the deltoid insertion on the

![Figure 7](https://example.com/figure7)

(A) A preoperative x-ray image. (B) A postoperative x-ray image shows placement of cerclage wires. (C) A preoperative x-ray. (D) A postoperative x-ray image shows placement of cerclage wires.
lateral side of the vertical osteotomy. The VHO technique leaves the distal cement plug intact, therefore avoiding the use of long-stem revision implants and the extensive bone loss or diaphyseal perforation that can follow the attempted removal. The proposed osteotomy approach is also quite expedient. The entire osteotomy and stem extraction requires only 10 to 15 minutes.

Revision shoulder arthroplasty presents a technical challenge with regard to implant removal and replacement. New prosthetic designs offer surgeons a wide variety of replacement implants; however, it is still essential to successfully remove the failed prosthesis without causing further bone loss or iatrogenic injury. Nevertheless, as recently described in the literature, revision shoulder arthroplasty can result in a significant improvement in outcome scores for the patient.\(^8\) When an arthroplasty fails, it is most likely at the glenoid through either loosening of the component or progression of arthrosis.\(^24,30,41\) The humeral component remains well-fixed in most cases. The proposed technique greatly facilitates the removal of the humeral stem in a safe and controlled manner.

**Conclusion**

In conclusion, although long-term follow-up studies are needed to evaluate the potential for postoperative peri-prosthetic fracture or humeral stem loosening, or both, with this technique, we have not seen these complications in our patient population. The vertical humeral unicortical osteotomy does facilitate efficient humeral stem removal, with no compromise of proximal bone, and can be recommended in revision shoulder arthroplasty cases.

**Disclaimer**

Dr Nicholson is a designer and consultant for Zimmer. Dr Romeo is a designer and consultant for Arthrex. The other authors, their immediate families, and any research foundations with which they are affiliated have not received any financial payments or other benefits from any commercial entity related to the subject of this article.

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