SHOULDER ARTHROPLASTY (G ATHWAL, SECTION EDITOR)

Platform systems in shoulder arthroplasty

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Abstract A platform system for shoulder arthroplasty can include a convertible modular humeral stem and/or a metal-backed glenoid component to facilitate straightforward conversion from either a hemiarthroplasty or anatomic total shoulder arthroplasty to a subsequent anatomic total shoulder arthroplasty, or more frequently, reverse total shoulder arthroplasty without any revision to the stem and/or glenoid baseplate. Recent studies have demonstrated a decreased rate of intraoperative humeral fracture, complications, and blood loss when a platform system is used and the humeral stem is not exchanged during revision arthroplasty. Future studies with larger patient cohorts are necessary to truly evaluate the potential value and limitations of this technology.

Keywords Platform system · Total shoulder arthroplasty · Reverse total shoulder arthroplasty · Shoulder hemiarthroplasty · Revision shoulder arthroplasty

Introduction

The volume of shoulder arthroplasty procedures performed in the USA has increased substantially over the past decade,

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from 14,000 shoulder hemiarthroplasty (HA) and total shoulder arthroplasty (TSA) procedures performed in 2000 to over 46,000 performed in 2008, an increase of nearly 12 % per year [1]. If this trend continues, an estimated 100,000 shoulder arthroplasties will be performed annually this year [2]. While at least part of these increases are due to an aging population that desires to remain active, a large percentage of this increase is due to rising popularity of and expanding indications for reverse total shoulder arthroplasty (RTSA) [3•].

Accompanying this exponential increase in primary shoulder arthroplasty is an expected increase in the incidence of revision shoulder arthroplasty. The estimated rate of revision for failed shoulder arthroplasties has grown by 400 % over the last 20 years, making revisions account for up to 10 % of all shoulder arthroplasty [4–6]. Revision shoulder arthroplasty is most often indicated for multiple modes of failure, with glenoid component loosening being the most common. Revision arthroplasty can be significantly more challenging than the index procedure due to the frequent need for component removal, which can lead to bone loss [7].

Recently, due to success of reverse total shoulder arthroplasty for rotator cuff tear arthropathy, the indications have expanded to use of this procedure as a revision option or salvage for failed prior arthroplasty, including revision of failed shoulder hemiarthroplasty or failed anatomic shoulder arthroplasty [8, 9•]. Conversion of prior arthroplasty to a reverse shoulder arthroplasty can be technically demanding and fraught with complications. These challenges led to the recent introduction of a platform system for shoulder arthroplasty, which can include a convertible modular humeral stem and/ or a metal-backed glenoid component to facilitate straightforward conversion from either a hemiarthroplasty or anatomic total shoulder arthroplasty (ATSA) to a RTSA without removal of the stem or glenoid baseplate [10•].



Uses and advantages of a platform system

A platform system for shoulder arthroplasty offers system flexibility and interchangeability for subsequent revision surgery. This would allow, in theory, for a less demanding and time-consuming revision surgery, and potentially reduce complications associated with the removal of glenoid or humeral components. A platform system consists of two potential modular components: a versatile humeral stem and a metal-backed glenoid component.

Humeral stem in a platform system for shoulder arthroplasty

Conversion of a traditional HA or ATSA to RTSA requires the removal and reimplantation of a humeral stem. As humeral stem loosening is exceedingly rare due to good integration with host bone, humeral shaft corticotomy is often required for stem removal, which is associated with a high rate of fracture [11, 12]. Additional fracture risk may rise from cerclage wire fixation which may be required to stabilize diaphyseal osteotomies or from diaphyseal perforation of the revision stem. One recent study noted a 21 % intraoperative humeral shaft or tuberosity fracture rate during stem removal during conversion to RTSA [13]. Other studies have noted humeral fracture rates of up to 25 % during stem removal at revision [10•, 12, 14, 15]. Additional risks of humeral stem removal include potential loss of humeral bone stock, nerve injury, periprosthetic fracture, and malunion or nonunion of a humeral osteotomy with later humeral component loosening [11, 14, 16, 17•].

The use of a platform system allows conversion of a hemiarthroplasty or ATSA to RTSA without removal of the humeral stem (Table 1). Compared to patients in whom stem removal is required, this virtually eliminates the risk of intraoperative humeral fracture. A recent study reported a 0 % rate of humeral tuberosity or shaft fractures in patients with platform humeral stems

Table 1 Current commercially available platform systems in the USA

Manufacturer/name	HA option?	ATSA option?	RTSA option?
Biomet Comprehensive	Yes	Yes	Yes
Depuy Global Unite	Yes	Yes	Yes
Exactech Equinoxe	Yes	Yes	Yes
Tornier Ascend Flex	Yes	Yes	Yes
Integra Titan	Yes	Yes	Yes
Smith & Nephew PROMOS	Yes	Yes	Yes
Lavender Medical UNIC	Yes	Yes	Yes

HA hemiarthroplasty, *ATSA* anatomic total shoulder arthroplasty, *RTSA* reverse total shoulder arthroplasty

converted to RTSA [13]. A similar study of 26 patients with full modular stems likewise reported no intraoperative humeral complications [18].

Metal-backed glenoid component in a platform system for shoulder arthroplasty

Glenoid component loosening is one of the most common complications in ATSA and represents 25 % of all complications related to TSA [19]. Often, glenoid component loosening is not isolated, but combined with a rotator cuff tear, glenohumeral instability or component malposition [20]. Glenoid revision is indicated for painful glenoid loosening or glenoid malposition or wear in instances where the rotator cuff remains competent. Often, as mentioned before, glenoid bone loss and rotator cuff deficiency, particularly the subscapularis, often necessitate conversion to a glenosphere for RTSA at the time of revision [10•]. A platform system utilizes a versatile metal-backed glenoid that allows simple exchange of the polyethylene component when revising ATSA, or conversion to a glenosphere when converting from ATSA to RTSA.

Metal-backed glenoids are not a new concept; they have been debated since Neer and colleagues published their results in the 1980s [21]. Numerous authors have published failures of metal-backed components. Taunton and co-authors reported on 83 ATSA with a metalbacked, bone-ingrowth glenoid component and found a 40 % rate of glenoid component loosening and 25 % rate of polyethylene wear with metal wear of the glenoid component [22]. Boileau, et al reported a 20 % loosening rate of metal-backed glenoid components compared to 0 % in all polyethylene glenoids in a prospective study of 40 patients undergoing ATSA [23]. Improvements in modern implant design may ameliorate these issues, as a recent study that included twelve primary platform ATSA with metal-backed glenoid components revised to RTSA did not report any metal-backed glenoid revisions [10•].

Clinical outcomes using platform systems

As platform systems are a relatively new advent in the evolution of shoulder arthroplasty design, only limited case series exist describing clinical outcomes utilizing such technology. Castagna, et al recently published a series of 26 patients who underwent conversion of a previous shoulder arthroplasty to an RTSA using a platform system [18]. The index operation in 18 of the patients was hemiarthroplasty for fracture; the remaining 8 patients had an ATSA for glenohumeral arthritis. The authors reported that the use of a fully modular system allowed avoidance of humeral stem removal in all

patients, and the metal-back glenoid in revision of ATSA, resulting in a short operative time and few intraoperative complications (Fig. 1). At an average of 32 months follow-up, the average constant score improved from 25 points to 48 points, while the EuroQol Visual Analog Scale (EQ-VAS) improved from 40 preoperatively to 70 postoperatively. All patients improved in terms of range of motion and follow-up imaging obtained at final follow-up demonstrated good integration of the implants without any evidence of loosening [18].

A recent study published by Wieser, et al evaluated outcomes in 48 patients with hemiarthroplasties and eight patients with ATSA converted to RTSA with (n=43) and without (n=13) stem exchange to elucidate potential advantages of a platform system [17•]. The authors found significantly shorter surgical time and lower average blood loss in patients without stem exchange. Intraoperative complications were also significantly lower in patients without stem exchange (8 %) compared to patients with stem exchange (30 %). Clinical outcomes were assessed in patients who did not have intraoperative complications. Average Constant scores improved from 30 to 48 in the stem-retaining group, which was not significantly different from the stem-removal group (in patients without intraoperative complications), who had an improvement from 24 to 45 [17•]. Although these results are promising, several limitations of this study should be noted. The average time to revision of the index shoulder arthroplasty was only 38 months, which is quite short and raises the question of component malposition as an etiology for many of the revisions included in the study. Furthermore, the authors state that improper placement of the initial stem may preclude retention; this idea is supported by the fact that one third of the shoulders underwent removal of a modular stem, and those patients who underwent exchange of a modular stem often required subsequent surgery [17•].

Kany, et al recently reported their experience with a platform shoulder arthroplasty system in 29 patients, including five hemiarthroplasties, eight ATSA with cemented glenoid components and 16 ATSA with metal-backed glenoid components [10•]. Of these, three ATSA with cemented glenoids were converted to ATSA with metal-backed glenoids, and the remainder of the included arthroplasties was converted to RTSA. The authors found that the humeral stem could be maintained in 72 % of patients, with the remaining patients requiring stem change due to a high position that prevented reduction of the new RTSA. Corticotomy or humeral shaft windows were not required for stem removal. Only two metal-backed glenoid components required exchange during conversion from ATSA to RTSA, both due to loosening in the setting of Walch type B2 glenoid bone wear. Clinically, patients demonstrated significant improvement, from an average preoperative Constant score of 27 to 60, postoperatively. At a mean follow-up of 28 months, no radiological glenoid or stem loosening was noted [10•].

Limitations and complications associated with platform systems

While designed to reduce intraoperative complications such as blood loss, humeral fracture and glenoid bone loss due to the time and effort involved in component removal during revision shoulder arthroplasty, platform systems have associated limitations and complications that require mention. When choosing to maintain a convertible humeral prosthetic stem, the surgeon must

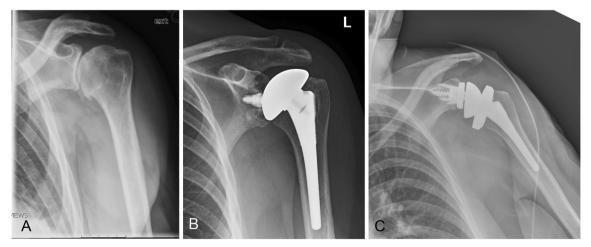


Fig. 1 A 72-year-old female with left shoulder pain due to osteoarthritis (**a**) underwent anatomic total shoulder arthroplasty using a convertible humeral stem (**b**). She later fell and suffered a subscapularis rupture,

which failed primary repair. She was later converted to an RTSA, maintaining her previous stem, with an excellent functional result (c)

accept the position of the stem, which is certainly a limitation if the position is suboptimal [9•]. Care must be taken at the index operation to utilize meticulous surgical technique to appropriately position the stem, not only to improve the longevity of the index implant, but to allow for revision without any need for stem removal. Based on existing studies, even when presumably good surgical technique is utilized, it is possible that removal of a convertible stem may be required at the time of revision to appropriately reduce the revision prosthesis and achieve adequate soft tissue tension in revision to RTSA. In addition to humeral length, excessive or abnormal humeral version may also require humeral component removal especially in revision to RTSA.

Limitations of a metal-backed glenoid component, as discussed previously, also exist. With improvements in design, wear characteristics have likely improved since the initial results published by Neer [21], although no clinical outcomes studies have confirmed this. For surgeons concerned with the implications of utilizing a metal-backed glenoid, systems exist in which only the humeral stem is convertible and a traditional glenoid component can be used.

Conclusions

Platform systems in shoulder arthroplasty offer significant versatility and numerous advantages, particularly in the revision setting, by decreasing the amount of surgical exposure needed during revision surgery, often eliminating the need for humeral osteotomy and thus reducing the incidence of intraoperative humeral fractures, decreasing surgical time, decreasing blood loss and decreasing overall complications during conversion from previous arthroplasty. Further studies with larger patient cohorts are necessary, however, as the existing literature is limited to small case series with heterogeneous populations.

Compliance with ethical standards

Conflict of interest Brian C. Werner declares that he has no conflict of interest.

Joshua S. Dines reports grants and personal fees from Arthrex, personal fees from Conmed Linvatec, and personal fees from Ossur, outside the submitted work.

David M. Dines reports royalties received from Biomet Inc. and consulting fees from Tornier Inc.. In addition he is the developer of the Biomet Comprehensive Reverse Total Shoulder System.

Human and animal rights and informed consent This article does not contain any studies with human or animal subjects performed by any of the authors.

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