Metallic Modular Taper Junctions in Total Hip Arthroplasty

McTighe T1, Brazil D2, Clarke I3, Keppler L4, Keggi J5, Tkach T6, McPherson E7

Abstract

The emergence of modularity in total hip arthroplasty (THA) in the 1980s and 1990s was based on the fact that the benefit of these design features outweighed the risk. The use of metallic modular junctions presents a unique set of advantages and problems for use in THA. The advantages include improvement in fit and fill of the implant to bone, restoration of joint mechanics, reduced complications in revision surgery and reduction of costly inventory. However, the risks or concerns are a little harder to identify and deal with. Certainly corrosion, and fatigue failure are the two most prevalent concerns but now the specifics of fretting wear and corrosive wear increasing particulate debris and the potential biological response is having an impact on the design and potential longevity of the reconstructed hip. Material and designs are facing a shorter life expectancy than what was previously thought, mostly due to an increasing level of physical activity by the patient. Because there are no accurate laboratory test whereby the service life and performance of these implants can be predicted, early controlled clinical evaluations are necessary. Early publication of testing and clinical impressions should be encouraged in an attempt to reduce exposure to potential at risk patients, implants and material. The reduction and possible elimination of risks will require a balancing of all the variables requiring a multidisciplinary endeavor.

This paper is designed to review the risk factors, and benefits of modular junctions in total hip arthroplasty (THA). Also some basic engineering principals that can reduce risk factors and improve functionality of modular junctions.

Keywords: hip, arthroplasty, debris, fretting, modularity, taper, metal ions, and metallurgy

Level of Evidence: AAOS Therapeutic Level III

Introduction

In dealing with the vast and complex problems associated with reconstructive total hip arthroplasty (THA), one of our tools is the use of metallic modular junctions. [1,2,3,4] Recently there has been considerable discussion and debate surrounding the risk benefit ratio is using modularity. [5,6,7] Modularity selected for THA is typically determined by ascertaining the intended function of the modular junction in the overall reconstruction of the hip. The most
suitable modular designs are those that are well tolerated by the body and can withstand increased cyclic loading in an ever-demanding environment, especially with the physical activities and expectations of today’s patients. Often, the totality of factors that must be assessed when choosing a modular junction for implantation is not completely considered. Typically, the surgeon considers only one issue, which is material strength. Other critical factors of modularity selection include corrosion resistance, cost, and ability to manufacture. [3,6,8]

Individual modular design parameters can offer significant advantages for both fit and fill of implant to bony structures while providing more options for intraoperative customization of joint mechanics and significant economic value in reducing levels of finished goods inventory (Figure 1). [6,9] Now, amid reports of clinical incidents in which metal modular junctions have demonstrated fretting, corrosion, and pseudotumors, there is renewed interest as to what causes these junctions to fail. [9,10,11,12,13] The recent fall in the use of modularity can be contributed primarily to concerns with inflammatory reactions to metal debris. Can failures be predicted or avoided? When a failure does occur what can be done about it?

**Current Concerns with Metallic Materials**

Implant compatibility and particulate debris in THA is not a new concern and has been an issue of debate since the first attempt to replace a hip joint in the 1890s. One of our greatest allies in reconstruction is the use of metals for implant fabrication; however, this requires an understanding of the biological and engineering principles involved. [8,14]

While modular designs represent an advance in the ability to precisely fit the implant to the bone and restore joint mechanics, the mechanical integrity of the assembled component must be fully tested before clinical use. Fabrication methods, tolerances, surface characteristics, materials, electrochemical environment and mechanical environment are all critical factors that need careful consideration in evaluating the long-term performance of modular interfaces. In evaluating the mechanical performance of modular femoral stems, there is no single test that can adequately represent the various conditions that a hip stem maybe subjected to in vivo.

Biocompatibility is mainly determined by the implant surface properties. When a metal implant comes in contact with biological tissue, the following occurs:

1. The implant is first covered with proteins from the body fluids, then cells may attach according to the implant surface properties.

2. The body will either tolerate a biocompatible implant or a foreign body reaction will occur. For metals, this depends on the surface properties of the implant, such as surface chemistry and roughness. Proteins and cells interact differently on surfaces with different properties (Figure 2). If the implant is biocompatible, the inflammation will decrease. If the implant is not biocompatible, a chronic inflammation can occur with possible consequence of a foreign body reaction. In addition, damaged surfaces may evolve to release ions that are potentially allergenic/toxic. This is the beginning of the corrosion process (Figure 3,4,5).

It seems that every 10 years, concern regarding problems from implanted materials resurface. It has been almost four decades since Willert first described the problem of polyethylene wear leading to peri-prosthetic inflammation, granuloma, bone resorption, and implant loosening. [15] Bobyn et al presented an AAOS scientific exhibit in 1993 reviewing problems and solutions with particulate debris in THA. [10] This review covered concerns with
modularity (tapers, dovetails, pads, and stem segments) in both the femoral and acetabular component. So what is different today? Why the increased concern?

Material selection and fabrication has not been altered to any great degree since the 1990s. However, three significant factors have come into play. First, volume of total joint surgery has increased (U.S.), and primary THA is projected to increase by 174% to 572,000 per year by 2030. Second, THA is being done on younger patients and patient activity overall within all age groups has increased. Third, small design alterations may have significant negative outcomes. [7,8,9,10,13]

Another possible factor is the reluctance of surgeons to provide postoperative precautions with regard to early physical activities. Regardless of material or design, the surgical process for preparing and inserting a total hip stem requires a fracture healing response of the bone. Bone remodeling initially occurs under the stable condition of fracture with rigid fixation and no gap formation—the key being stability of implant to bone to maintain the biological healing response. [17] Modular junctions are designed to work in a stable environment. If the implant has instability and micro-motion, it is very likely the modular junction will encounter increased stress that can lead to a breakdown of the stability of the modular junction, which results in fretting and or corrosion.

Recent concerns with modular tapers can be attributed to the results with metal-on-metal (M-o-M) hip resurfacing (HR) and by extension, the use of large heads (greater than 36 mm) in THA. [18] Small diameter heads (28-32 mm) have had favorable results since the late 1980s. [19,20,21] However, the market demand to reduce dislocations in THA pushed the M-o-M bearings into larger head diameters. While it took time to see the problems with large M-o-M heads, it is also possible that the signs were overlooked. Since 1956, there have been reports of soft tissue tumors caused by metallic alloys. [22] By 1998, Jacobs reported that the taper junction between head and stem was responsible for the significant increase in titanium and cobalt concentrations in the patient, even when the prostheses were
functioning well [23] (Figure 6). In the 2010 National Joint Registries of England and Wales (NJR), problems were becoming obvious with the focus being directed to the taper junction. [24] In 2012, the Medicines and Healthcare products Regulatory Agency (MHRA) in the United Kingdom (UK) issued new guidelines on larger head (+36mm) forms of “M-o-M” hip implants. Patients with a large M-o-M hip implant should have annual health checks for life as compared to previous recommendation of up to five years. (Figure 7a, 7b, 7c, 7d).

In May 2015, Michael Morlock published a review paper on tapers showing examples of head/neck taper fractures with a Ti-alloy stem taper and a titanium sleeve connector to the femoral head [25] (Figure 8). He further...
pointed out in his paper that the European Union with the establishment of a Scientific Committee on Emerging and Newly Identified Health Risks (SCENIHR) working group investigating “The safety of M-o-M joint replacements with a particular focus on hip implants about the M-o-M problems.”

The preliminary consensus of this working group was published in September 2014 and addressed this topic explicitly: “This metal debris can originate either from the bearing articulation directly or from the modular taper junction between prosthesis head and stem. In the past, the taper has only been reported anecdotally as the origin of problems. Recently, the taper has emerged as the focus of attention, since large modular metal heads for M-o-M arthroplasty were introduced due to their ability to reduce dislocation risk, which is the second major complication in hip arthroplasty. These larger heads, however, put larger loads on the taper junction and are suspected to be responsible for the problems suddenly occurring at this side.” [21,26]

The success of a self-locking taper is influenced by the design of the taper, particularly the taper angle, the roughness, and the mating materials between the “male” and “female” components. (Figure 9a, 9b) This results in co-integration (locking), with material transfer across the zone of contact (cold welds). The degree of fit (interference) is determined by the relative dimensions of the two components (male and female regions), and a design decision to have interference along a specific part of the taper’s circumference and length. The area of interference contact must be adequate to maintain integrity under functional (loaded) conditions, while the surface finish of the components must be specific to the physical and mechanical properties of each component’s material. [17,34]

In the last two decades, manufactures have been altering femoral stem trunnions from various tapers such as 14/16 to 12/14 to 11/13 (Figure 10). The Ceramtec 12/14 taper at one time has been referred to by most in Europe as a European 12/14 taper. This term was not trademarked, and some companies began altering the manufacturing tolerance as originally produced from Ceramtec. The term “Euro taper” still is used by most in Europe to describe an off-the-shelf 12/14 Ceramtec taper. [27,28]

A range of different Morse taper angles, component tolerances and sizes, and surface finishes exist within commercially available hip systems. While manufacturers do not recommend mixing and matching of component brands, a number of surgeons have been mixing and matching without complications, provided the products used have the same manufacturing tolerances. [29] A survey published in 2005 from the New Zealand Orthopaedic Association showed that 23% of the surgeons had implanted mismatched components within the last five years [36] (Figure 11).
Amid rising concerns of modular junctions, it is important to remember most hip implant revisions are not the cause of modularity. Aseptic loosening, osteolysis/wear, instability/dislocation, infection and periprosthetic fracture remain as the major reasons for hip revision surgery. One reason for revision that is growing in frequency is the failure of large M-o-M bearings. [30]

It is important to remember that early introduction of stem modularity did present problems, including disassociation of modular heads, incorrect head diameters implanted, and trunnion fatigue fractures [31] (Figure 12a, 12b). Unique to modular head/neck designs is the risk of dissociation of the head in association with dislocation and attempts at closed reduction. [43] This can often leave a well-fixed stem in place with some degree of damage to the stem trunnion. This may be an indication to use a modular trunnion sleeve to engage with the modular head, especially if you intend to use a ceramic head (Figure 13). Multi-modularity in stem designs, along with the use of larger head diameters, brings with it serious concerns with regard to corrosion and its biological reaction to increased metal ions and particulate debris.

**Corrosion**

Corrosion of metals has many different mechanisms that all have independent driving forces. Corrosion can be defined as the degradation of a material due to a reaction with its environment. There are many forms of corrosion and no universally accepted terminology is in use. The following terminology is based on current use by NASA-Kennedy Space Center. [32]

**Galvanic corrosion**

Galvanic corrosion is an electrochemical action of two dissimilar metals in the presence of an electrolyte and an electron conductive path. It occurs when dissimilar metals are in contact.

Worldwide ISO standards recognize the detrimental effect of galvanic corrosion cells that can be established in the body, and this should be considered during implant design. [8] When reduced taper length is combined with larger femoral heads, the outcome has been that industry experiences a new failure mode in THA “trunnionosis.” [33] One factor that can drive the trunnionosis phenomena is the use of different materials at modular junctions. Fundamental science states that two different materials in a conducting media will generate a battery or corrosion cell. Consequently, all differing materials mated together in the human body will set up a corrosion cell to some extent. The extent on the corrosion cell is affected by the fluid conductivity and galvanic potential difference between the two materials. [8]

**Pitting corrosion**

Pitting corrosion is localized corrosion that occurs at microscopic defects on a metal surface. The pits are often found underneath surface deposits caused by corrosion product accumulation.

**Crevice corrosion**

Crevic or contact corrosion is the corrosion produced at the region of contact of metals with metals or metals with nonmetals.

**Stress corrosion**

Stress corrosion cracking is caused by the simultaneous effects of tensile stress and a specific corrosive environment. Stresses may be due to applied loads, residual stresses from the manufacturing process, or a combination of both.

**Corrosion fatigue**

Corrosion fatigue is a special case of stress corrosion
caused by the combined effects of cyclic stress and corrosion. No metal is immune from some reduction of its resistance to cyclic stressing if the metal is in a corrosive environment.

**Fretting corrosion**

Fretting corrosion is the rapid corrosion that occurs at the interface between contacting, highly loaded metal surfaces when subjected to slight vibratory motions (Figure 14a, 14b).

Using these definitions, one can better understand the mechanisms behind product deterioration among the different THA junctions.

**Challenges with the Neck/Stem Modular Junction**

Fretting corrosion has recently been attributed to the decline in the clinical acceptance of modular neck hip implants. It has also been the reason for the recall of two products (Rejuvanate™ and ABGII™) by Stryker Orthopaedics, Mahwah, NJ. [7,8,34,35,36] A main driving mechanism behind fretting corrosion is stress, or load. Increasing the stress at the modular junction will proportionally increase the extent of the fretting corrosion (Figure 15). Reviewing the design of the modular junction of these products indicates that the application of some fundamental engineering principles could have reduced the probability of fretting corrosion.

Figure 14a. Schematic illustrating that oscillatory motions as small as 10-100µm can induce wear and mechanically-assisted crevice corrosion (MACC). Such relative motion is unavoidable because modular junctions inherently introduce parts of different rigidity (size, shape) and different alloys (stiffness criteria). (Courtesy of Ian Clarke)

Figure 15. Retrieval of taper corrosion with dissimilar metals—cobalt-chrome alloy modular neck on titanium Stryker stem. (Courtesy of Dartmouth Biomedical Engineering Center.)

Figure 14b. Electrochemical reactions involved in mechanically-assisted crevice corrosion (MACC: redrawn from data in Gilbert et al., 1997). Cyclic loading is a necessary hip function and the resulting micromotion can disrupt the protective oxide film on metal surfaces. Ideally this will quickly reform, the metal combining with oxygen from the local environment and in the process releasing hydrogen and negatively-charged electrons. Under adverse conditions, the fretting continues, the oxygen concentration is depleted, and the formation of a protective oxide layer is compromised. In addition the surrounding environment is becoming more acidic as the hydrogen ions recombine with chloride ions to form hydrochloric acid, thereby promoting dissolution of the metal surface. (Gilbert et al 1997) (Courtesy of Ian Clarke)
potentially leads to a more rapid fretting corrosion rate as compared with neck preserving style stems (Figure 16d). [7,8] Figure 17 shows results of presentation by Brazil and McTighe on FEA modeling comparing level of neck resection (neck sparing stem versus conventional); as compared with a conventional neck resection, the neck sparing resection results in a 35% reduction in principal tensile stress. [46]

Recent marketing trends have also contributed to problems at the modular junction. The use of large femoral heads (greater than 36 mm) M-o-M bearings, increased femoral offset, increased leg length, and reduced precautions on patient-related physical activity may result in higher stresses at the modular junction. [7,8] These actions increase torque moment at the modular implant interface. On average, a 1-mm true lateral increase to the ball center offset will increase torque values by 8%. A 1-mm increase in vertical height (leg length) will increase torque by 6% (Figure 18).

Reduced taper engagement area, along with increased patient body weight and increased physical activity levels, places significant torsional loads on the implant. Torque
is a force applied over a distance (lever arm) that causes rotation about a fulcrum (axis of rotation) (Torque=Force (Fm) x Moment Arm). The greater the torque a muscle can produce, the greater the movement it will produce on the body’s levers. [45] Example of patient at risk would be an active male weighing 250 lbs with a 50 mm femoral offset, a combination that would generate in excess of 70 ft-lbs of torque. Design limit for most tapers is approximately 60 ft-lbs. We know by previous reports that the hip sees torque values over 95 ft-lbs, as demonstrated in some mechanical failures of first generation modular hip stems [13] (Figure 19).

One such torsional failure mode was presented as a poster exhibit at the 2006 ISTA Annual Meeting reporting on a proximal modular neck design that featured a “Dual Press™” modular junction. The Dual Press modular junction employs two areas of cylindrical press-fit (Figure 20). This allows the proximal portion of the shoulder to fully seat, providing medial support, which increases strength and allows higher lateral offsets. The rotation of the proximal body is restricted by a locating pin. The pin strength was established at 95 ft-lbs, well above historical published reports on torsion. These modular junction failures were not a fatigue failure mode, and no surgical errors or fabrication defects were found. The culprit appeared to be patient activity resulting in a mechanical overload in a static shear mode failure (perfect storm). The solution was rather simple: replace the old pin diameter from .125” to .188” and change the old plug to a new feature of a bolt that engages the stem. This revision resulted in 225% increase in torsional strength. It serves as an example that changes and improvements are possible once there is a full understanding of the problem. There have been no reported mechanical failures of its modular junction since 2004 with the improved design (Figure 21a, 21b, 21c).

Since 2004, there have been more than 7,000 Omni

![Figure 20. Dual-Press Modular Junction (Omni, East Taunton, MA) Illustration Showing two areas of press fit allowing proximal shoulder to sit flush with stem body.](image)

![Figure 21. Chart Showing Torque Loads Generated by Femoral Offset and Body Weight.](image)

![Figure 21a. Illustration showing old Dual-Press design to new improved design increasing torsional resistance from 95 ft-lbs to 216 ft-lbs.](image)

![Figure 21b. Explanted Apex Modular Stem (Dual Press Modular Junction, Omni, East Taunton, MA) showing sheared de-rotation pin and fretting abrasion wear. No signs of corrosion. (Courtesy of Keggi) (image)

![Figure 21c. Picture showing old pin diameter of .125” to new diameter of .188” increase in strength (+225%). (Courtesy of K. Keggi) (image)
MOD II and more than 3,000 Omni K2 Dual Press improved junctions implanted. Seventeen revisions involving the OMNI MOD (.23%) and four involving the OMNI K2 Stems (0.12%) have been reported to OMNI. Of these, two involved increased metal ion levels (as determined by the patient’s physician), and in both cases the OMNI MOD Stem was used with another manufacturer’s M-o-M femoral head and acetabular cup bearing combination. The revisions involved removing the competitor’s head and cup and replacing the OMNI MOD Modular Neck, leaving the stem in place. There have been no other reports of metal ion concerns, corrosion, or fretting with the OMNI MOD and OMNI K2 Modular Stems. [47]

Another example of modular neck failure was the original OTI™ Co-Cr-Mo modular neck that interfaced with a Co-Cr-Mo stem. The failure mode for this device was basic fatigue failure caused by an under-designed modular junction. Improvements made to this novel neck design, which increased surface contact by 40%, included specific size increases of the taper trunnion that improved mechanical strength from 520-700 lbs to greater than 1,200 lbs (Figure 22a and 22b). To our knowledge, there has been no reported failures with the improved modular junction design. [48,49]

Our own research using a short-curved neck sparing modular neck that mates with a Co-Cr-Mo neck with Ti-alloy stem is undergoing extensive fretting and corrosion testing of the additional surface coating of selected regions of the Co-Cr-Mo modular neck with titanium nitride (TiN). This material process reduces the potential galvanic reaction between the materials and consequently reduces the probability of corrosion between mating surfaces. This fundamental design concept can be further applied to the internal surfaces of the femoral head that interfaces with a Ti-alloy stem taper. Our initial results have been presented at various CME meetings with very favorable results in reduction of fretting abrasion wear between the TiN distal coated necks versus non coated necks. [7,8,36,37] TiN fully coated necks saw the same results for the distal coated necks but saw no difference interfacing with Co-Cr-Mo heads.

**Challenges with the Stem/Sleeve Modular Junction**

The success of the S-Rom® modular stem system stimulated most companies to rush into the market with a modular style hip. [9,38,39] The S-Rom stem, an evolution of the original Sivash stem, experienced a number of design changes before becoming the novel design that still survives today. Most think the modular features were the single most important factors to its success. In reality, the clinical success can be contributed to its basic geometric design that provided for immediate implant stability with the potential for long-term fixation with a reproducible surgical technique. The modular features are secondary to its basic geometric structure.

Fracture of the S-Rom stem is rare; however, it does happen, and fractures have been reported at different sites in the femoral stem body. Pearce et al reported two stem fractures at the mid-stem junction at the top of the slotted portion of the stem. [40,41,42] One of our authors had a fractured stem (4 years postoperatively) within the sleeve/
ethylene deposits. In 1992, Buly et al reported on 71 cases of titanium wear debris in failed cemented THA. Femoral bone loss in aseptically loose, primary THA was graded as severe in 51%, moderate in 24%, and mild in 20%. Femoral endosteolysis was present in 94%, while acetabular osteolysis was seen in 6%. Histological evaluation of tissues from failed primary arthroplasties revealed poly(methyl methacrylate) debris in 75% of cases, polyethylene debris in 80%, metal debris in 75%, and chronic inflammatory cells in all cases. One can conclude from past reports and personal observations that rougher surfaces that interface with another rough surface (bone, cement, metal) under micro or macro movement will suffer fretting abrasion wear.

Many manufactures use the bead blasted or matte finish on titanium stems as a cosmetic process to cover or reduce machine marks from the fabrication process.

One area of observation is the process of grit-blasting titanium stem surfaces, which leaves a matte or satin finish (Figure 25). Grit surfaces were introduced in the early 1980s for cemented stems, with the belief that the slightly rough surface finish (RA: .7µm) would provide improved bonding of the bone cement interface. Results were just the opposite, with higher aseptic loosening occurring in the grit-blasted stems than in polished stems. In retrieving roughened Spectron EF stems (Smith and Nephew, Memphis, TN, USA), Gross et al reported the presence of macroscopic metallosis in all hips. The microscopic examination of the femoral pseudomembrane consistently revealed an inflammatory reaction characterized by the presence of multinucleated giant cells and metallic, cement, and polyethylene deposits. In 1992, Buly et al reported on 71 cases of titanium wear debris in failed cemented THA. Femoral bone loss in aseptically loose, primary THA was graded as severe in 51%, moderate in 24%, and mild in 20%. Femoral endosteolysis was present in 94%, while acetabular osteolysis was seen in 6%. Histological evaluation of tissues from failed primary arthroplasties revealed poly(methyl methacrylate) debris in 75% of cases, polyethylene debris in 80%, metal debris in 75%, and chronic inflammatory cells in all cases. One can conclude from past reports and personal observations that rougher surfaces that interface with another rough surface (bone, cement, metal) under micro or macro movement will suffer fretting abrasion wear.

Many manufactures use the bead blasted or matte finish on titanium stems as a cosmetic process to cover or reduce machine marks from the fabrication process.

One area of observation is the process of grit-blasting titanium stem surfaces, which leaves a matte or satin finish (Figure 25). Grit surfaces were introduced in the early 1980s for cemented stems, with the belief that the slightly rough surface finish (RA: .7µm) would provide improved bonding of the bone cement interface. Results were just the opposite, with higher aseptic loosening occurring in the grit-blasted stems than in polished stems. In retrieving roughened Spectron EF stems (Smith and Nephew, Memphis, TN, USA), Gross et al reported the presence of macroscopic metallosis in all hips. The microscopic examination of the femoral pseudomembrane consistently revealed an inflammatory reaction characterized by the presence of multinucleated giant cells and metallic, cement, and polyethylene deposits. In 1992, Buly et al reported on 71 cases of titanium wear debris in failed cemented THA. Femoral bone loss in aseptically loose, primary THA was graded as severe in 51%, moderate in 24%, and mild in 20%. Femoral endosteolysis was present in 94%, while acetabular osteolysis was seen in 6%. Histological evaluation of tissues from failed primary arthroplasties revealed poly(methyl methacrylate) debris in 75% of cases, polyethylene debris in 80%, metal debris in 75%, and chronic inflammatory cells in all cases. One can conclude from past reports and personal observations that rougher surfaces that interface with another rough surface (bone, cement, metal) under micro or macro movement will suffer fretting abrasion wear.

Many manufactures use the bead blasted or matte finish on titanium stems as a cosmetic process to cover or reduce machine marks from the fabrication process.

One area of observation is the process of grit-blasting titanium stem surfaces, which leaves a matte or satin finish (Figure 25). Grit surfaces were introduced in the early 1980s for cemented stems, with the belief that the slightly rough surface finish (RA: .7µm) would provide improved bonding of the bone cement interface. Results were just the opposite, with higher aseptic loosening occurring in the grit-blasted stems than in polished stems. In retrieving roughened Spectron EF stems (Smith and Nephew, Memphis, TN, USA), Gross et al reported the presence of macroscopic metallosis in all hips. The microscopic examination of the femoral pseudomembrane consistently revealed an inflammatory reaction characterized by the presence of multinucleated giant cells and metallic, cement, and polyethylene deposits. In 1992, Buly et al reported on 71 cases of titanium wear debris in failed cemented THA. Femoral bone loss in aseptically loose, primary THA was graded as severe in 51%, moderate in 24%, and mild in 20%. Femoral endosteolysis was present in 94%, while acetabular osteolysis was seen in 6%. Histological evaluation of tissues from failed primary arthroplasties revealed poly(methyl methacrylate) debris in 75% of cases, polyethylene debris in 80%, metal debris in 75%, and chronic inflammatory cells in all cases. One can conclude from past reports and personal observations that rougher surfaces that interface with another rough surface (bone, cement, metal) under micro or macro movement will suffer fretting abrasion wear.

Many manufactures use the bead blasted or matte finish on titanium stems as a cosmetic process to cover or reduce machine marks from the fabrication process.
The original S-Rom 125° stem (1984) had a locking pin through the stem that engaged with the proximal sleeve, reducing the risk of stem slippage within the sleeve (Figure 27a and 27b). One additional problem with the 1984 design was a groove that ran down the anterior/posterior portion of the entire length of the stem. In a poster exhibit at the 2006 ISTA annual meeting, one of our coauthors presented an example of progressive distal osteolysis, in which particulate debris migrated down the grooved stem. This helped make the decision of adding distal flutes, eliminating the groove, and eliminating the locking pin. Another concern with regard to the concept of stem/sleeve modularity is the risk of increasing the length beyond the taper engagement contact zone. During the 1980s, one of our coauthors was part of the S-Rom design team; their study group (Cameron, Mallory, Bierbaum, Bobyn, Moorland, Pugh, Greenwald, Noiles and McTighe) reviewed the design and ruled it out because of increased risk of fretting abrasion wear between distal sleeve and stem (Figure 28). Bending moments can be increased, especially with thinner stem diameters.

Challenges with Head/Neck Modularity

As taper lengths and taper ratios have changed over the years, standardizing on a 12/14 Euro Ceramtec off-the-shelf style taper allows for more standard revision options as compared with using a taper neck sleeve adapter. Neck taper adapters may have limitations in design by having skirts that may interfere with range of motion or cause impingement, resulting in generation of particulate debris and/or dislocation (Figure 29).

This design concept allows for adjustment of head center vertical height and head center lateral offset. In addition, this feature allows for mixed materials to be selected for the articulation of the bearing surface. The femoral head is commonly fabricated from a Co-Cr-Mo alloy or an alumina-based ceramic. Our research on TiN of Co-Cr-Mo modular necks interfacing with Ti-alloys stems might be carried over to just coating the inside of a Co-Cr-Mo head to reduce potential galvanic reaction of dissimilar materials and reduction of micro-fretting abrasion wear at the head/neck interface. Another improvement already in practice is going back to the concept of a more hemispherical head. This improves the surface contact area for head/neck trunnions that can reduce stress and micro-motion at the interface (Figure 30).

Summary and Conclusion

The use of metallic modular junctions in hip replacement has increased since the early 1980s, and some might say they are overused. We are seeing an increased num-
ber of complications associated with modularity, including dissociation, corrosion, wear, fretting, and fatigue failure. When modular implants were first introduced, the biggest challenge was the frequent fracture of ceramic heads. Today—more than 40 years after the introduction of modular ceramic heads—fracture is rare, and ceramic modular heads have demonstrated low wear rates, outstanding biocompatibility, diamond-like hardness, and high resistance to third-body wear.

Metallic heads made of titanium alloy proved to be unsatisfactory, increasing wear at the articulation. Metallic heads made from Co-Cr-Mo alloy are strong, and they pose no potential of failure by fracture or fatigue. However, fretting, corrosion, and micro-motion are still major concerns. Modular implants with titanium alloy stems and Co-Cr alloy heads were introduced in order to take advantage of the lower stiffness of Ti alloy for better load transfer to bone while making use of higher wear resistance of Co-Cr alloy heads. The use of these modular implants soon gave rise to corrosion at the modular mating surfaces, which was first thought to be galvanic in nature because the dissimilar materials were involved. Further investigations on Morse taper connections of modular hip prostheses brought about different conclusions on the nature of modular interface failures. Stress, strain, and micromotion at the modular interface can induce fretting abrasion wear, resulting in the generation of particulate debris, increased release of metal ions, corrosion, and adverse tissue (local and systemic) reaction.

Analysis shows that you must carefully consider patient weight and activity level when implanting a hip stem of any design. A 350-pound active male with a 50 mm offset and a 11 mm distal stem exceeds the fundamental fatigue strength of titanium alloy, regardless of proximal stem design or modularity.

Taper issues are the same regardless of where the taper is located (head/neck, neck/stem, mid-stem, or stem/sleeve). Reducing risk associated with tapers can be ameliorated through many strategies; design characteristics such as a large surface contact (length and diameter), stiffer material (less deflection), and tight manufacturing tolerances can reduce stress, strain and micro-motion at the modular junction. This reduces fluid ingress and the extent of fretting that could trigger corrosion by depassivating the protective metallic oxide layers and setting up a crevice corrosion cell. Careful intraoperative techniques for assembly are critical. Both male and female trunnions must be clean and dry before assembly, and proper force must be used to engage the modular junction.

Generation of particulate debris can often be reduced through the careful selection of implant material and fabrication. This problem is worse with Ti-based implants because of lower hardness and abrasion resistance. Also, some implant preparation techniques such as bead blasting tend to leave residual contaminants (silica or alumina) that can be dislodged by abrasion at the modular interface. Bead-blasted taper surfaces can produce surface scarring that is material transfer brought on by micro-motion. Taper surfaces should be clean and smooth, then micro-etched with chemical-milling techniques in the fabrication process. Debris can migrate throughout the joint space, accessing any and all implant interfaces. Select designs and material that provide immediate secure fixation that minimize micro-motion, stress, and strain.

The following are some examples of actions to reduce the generation of particulate debris:

- **Head/neck tapers:** Use 12/14 (Larger and stiffer surface contact area) taper over smaller tapers such as 11/13 or 9/10 when possible.
- **Head/neck tapers:** Increasing taper length will reduce micro-motion.
- **Stem tapers:** Many tapers do not have adequate intrinsic stability for high activity, so limit modular junctions or pick designs that have back-up features to support taper junctions (e.g., fluted stems).
- **Reduce fatigue failure of modular necks by material choice.** Co-Cr-Mo is stronger than Ti-alloy.
- **Reduce potential galvanic corrosion of dissimilar metals by TiN coating Co-Cr-Mo necks used with titanium stems.**
- **Reduce micro-motion, stress, and strain in modular necks by increasing taper engagement.**
- **Reduce micro-motion, torsional moment, and bending moment on stems (modular necks) by selecting neck sparing stem designs that retain the femoral neck.**
- **Caution should be used in selection of modular junctions in highly active males that exceeds 250 pounds.**

Modular designs have made significant contributions to reconstruction of the diseased and damaged hip—from improving fit and fill of the implants to restoring joint mechanics. While problems have been reported with the use of modularity, the collaborative orthopaedic community (industry, surgeons, and scientists) has been successful in identifying and providing solutions to improve overall designs and outcomes. Modularity can be designed and fabricated to provide safe, reliable, and reproducible clinical results.

As an example of industry stepping up identifying problems and initiating actions from 2002 to 2013, the six largest implant companies have voluntary recalled 578 hip implants as compared with the FDA using its recall authority three times in 20 years.
It is important to remember all devices are subject to failure. It is also necessary to recognize design and material limits and not to over-indicate in high-risk patients. A number of modular junctions have come and gone from clinical use. Nevertheless, the endeavor to improve clinical outcomes should be continued. Modularity can be designed and fabricated to provide safe, reliable, and reproducible clinical results.

Because there are no laboratory tests allowing accurate prediction of the service life and performance of implant parts, clinical experience with a large number of cases over a period of several years is the only reliable indicator. However, clinical evaluations should only begin after conducting aggressive basic science material and mechanical testing to anticipate potential failure modes. Individual patient physical activities should be considered when deciding on stem modularity features. Since there are no standards established for modular junctions the overall performance of modular junctions are not equal. Careful review of basic engineering principles is necessary and recognizing design limits will reduce the indication of overuse.

To advance scientific knowledge in the long run often requires some short-term setbacks.

Disclosure Statement:
One or more of our authors have disclosed information that may present potential for conflict of interest with this work. For full disclosures refer to last page of this journal.

Acknowledgement
The authors thank Chris Burgess, I. Clarke, A. John, J. Keggi, K. Keggi, Evert Smith, and Dartmouth Biomedical Engineering Center for pictures and illustrations.

References