

Early Results of a Modular Revision System in Total Knee Arthroplasty

Michael J. Morris, MD[†]; Keith R. Berend, MD[†]; Joanne B. Adams, BFA[†]; Adolph V. Lombardi, Jr., MD, FACS[†]

Abstract

Background: The number of revision total knee arthroplasty (TKA) procedures is projected to rise dramatically over the next fifteen years. These procedures are technically more demanding than routine primary TKA. Modular component options can assist the surgeon in addressing complex reconstructions in TKA, providing customization to remedy bony deficits, deformity, malalignment and instability. We review the early clinical results of a modular revision system that offers full interchangeability enhanced with a wide array of options for augmentation, offset, and constraint as well as modular stems.

Materials and Methods: A query of our practice's arthroplasty registry revealed a consecutive series of 100 patients (101 knees) who signed an IRB-approved general research consent allowing retrospective review, and underwent total knee arthroplasty performed with the modular revision system between May 2011 and May 2012. Reconstruction constructs and components, demographic variables, pre and post-operative clinical variables, failure modes, and survivorship were analyzed.

Results: At an average follow-up of 1 year (range, 0.1 to 2.2 years) there has been one aseptic failure for instability. One patient required incision and debridement of a non-healing wound. Three patients, all status-post reimplantation, failed secondary to recurrent infection, with one treated single-staged and the others undergoing another 2-staged exchange. Average ROM improved from 96.4° preoperatively to 104.8° at most recent evaluation. Average Knee Society clinical scores (0 to 100 possible) improved from 45.5 to 78.0, and function scores (0 to 100 possible) improved from 41.3 to 48.3. There have been no patient deaths during the follow-up period. Postoperative radiographs were available for review for 97 knees. Satisfactory position, fixation and alignment were observed in 92 (95%). Heterotopic ossification was observed around the tibial component of one knee. Minor radiolucencies were observed in femoral zones I and II and tibial zone 1 on lateral view of one knee, in tibial zones III and IV on AP view in one knee, and in tibial zone IV on AP view in one knee.

Conclusion: The early results of this modular TKA revision system are promising for use in complex TKA, with only one aseptic failure observed. There has been substantial improvements in ROM and function in this cohort.

Key words: Arthroplasty, Knee, Revision

† Joint Implant Surgeons, Inc.; Mount Carmel Health Systems New Albany, OH, USA

Introduction

Total knee arthroplasty (TKA) is a common procedure with great clinical success and 95% survivorship at 15 years as reported by multiple authors [1,2]. Annual rates of primary TKA are increasing, and projections for revision TKA are expected to rise 601% by the year 2030 [3]. There are significant technical challenges with revision TKA, including bone loss, deformity, malalignment, and instability coupled with a higher rate of both surgical and medical complications. The aim of this study was to investigate the early results of a fully interchangeable, modular revision knee system with a wide array of implant system options for sizing, stems, augmentation, offset, and constraint, designed to address the technical demands of revision surgery.

Materials and Methods

A retrospective investigation of our practice's arthroplasty database registry identified a consecutive series of 100 patients (101 knees) between May 2011 and May 2012 who underwent revision TKA with a modular revision knee system (Vanguard SSK 360, Biomet, Inc., Warsaw, IN, USA). A signed institutional review board (IRB) approved general research consent allowing retrospective review was obtained from all patients. The database was reviewed to analyze component constructs and level of constraint, demographic variables and preoperative and postoperative clinical assessment data including Knee Society scores, range of motion, limb alignment, indication for revision, and survivorship. Postoperative radiographs were reviewed in 97 knees.

There were 41 men and 59 women. Age averaged 64.2 years (range, 33-90) and BMI averaged 34.7 kg.m2 (range, 20-67). Procedures were revision of unicompartmental to total in 3 knees, revision in 69 (68%), and reimplantation after 2-stage treatment of infection in 29 (29%). Non-infectious indications for revision included aseptic loosening in 30 knees, instability in 38 knees, arthrofibrosis in 10 knees, periprosthetic fracture in 2, and malalignment in 2.

Two knees had no revision performed on the femoral side while 3 were revised to a cruciate retaining femoral component mated to the modular revision tibial component. Seven knees had no revision of the tibial component, while one was revised to a primary fixed I-beam tibial component mated with the modular revision femoral. The tibial inserts utilized were standard cruciate-retaining (CR) in one, CR-lipped in one, ultracongruent anterior stabilized in 3, nonconstrained posterior-stabilized in 12 and varus-valgus constrained in 84 (84%). Offset adapters were used with 42 femoral components and 28 tibial components. Average femoral and tibial offset, when utilized, was 4.8 and 4.7mm, respectively. A cruciate wing was added to 19 tibial components. Femoral and tibial stems were utilized in 96 and 91 knees, respectively. Femoral stems were 97% splined and with lengths of 40mm (n=6), 80mm (n=37), 120mm (n=45), 160mm (n=6), and 200mm (n=2). Tibial stems were 94% splined and with lengths of 40mm (n=3), 80mm (n=32), 120mm (n=56), and 160mm (n=3). Femoral augments were utilized in 52 cases and porous metal femoral augments in an additional 9 cases. Tibial augments were used in 26 cases and porous metal tibial augments in an additional 9 cases.

Results

Clinical outcomes improved significantly after the revision TKA compared with preoperative levels (Figure 1). At an average follow-up of 1 year (range, 0.1 to 2.2 years), range of motion improved from 96 degrees preoperatively to 105 degrees postoperatively. Knee Society clinical scores (range 0 to 100 possible) improved from 45.4 to 78.0, and function scores (0 to 100 possible) improved from 41.2 to 48.4.

One patient required incision and debridement of a non-healing wound. Three patients, all status-post reimplantation, failed secondary to recurrent infection, with one treated single-staged and the others undergoing another 2-stage treatment. One patient required polyethylene exchange for instability (polyethylene thickness increased 4mm). There have been no patient deaths during the follow-up period.

Postoperative radiographs were available for review for 98 knees. Satisfactory position, fixation and alignment was observed in 93 (95%). Heterotopic ossification was observed at the lateral tibial aspect in 1 knee with CR lipped bearing. A stable radiolucency was observed around the tibial component of one knee. Minor radiolucencies were observed in femoral zones I and II and tibial zone 1 on lateral view of one knee, in tibial zones III and IV on AP view in one knee, and in tibial zone IV on AP view in one knee.



Figure 1. A 68-year-old female patient presented with moderate left knee pain and instability 10 years after undergoing primary total knee arthroplasty with a cemented, posterior-stabilized device. Preoperative anteroposterior (A) and lateral (B) radiographs demonstrate aseptic loosening and tibiofemoral alignment of 8° varus. Anteroposterior (C) and lateral (D) radiographs at 2 years postoperative demonstrate well-fixed components in satisfactory position and alignment. Patient has been treated with left cemented revision TKA using a 62.5mm component with 10mm posterior augment, 2.5mm offset adaptor and 16x80mm splined stem on the femoral side, and 67mm tray with 16x63/67mm constrained insert, 16x80 splined stem, and small cruciate wing on the tibial side. The patellar component was not revised. Her Knee Society clinical score is 99 and function score is 50.

Discussion

The revision burden for TKA is increasing annually. Unfortunately, failure of revision TKA is not uncommon with some authors reporting rates as high as 63% within the first 5 years, predominately due to infection, instability, loosening, and patellofemoral problems [4]. These etiologies are also common in late failures, although polyethylene wear and aseptic loosening of cemented components predominates [5]. Additional risk factors for failure include younger age at the time of index arthroplasty, coronal malalignment, elevated body mass index, and lower socioeconomic and educational status [6]. Despite the various failure modes, the goal of revision arthroplasty, similarly to primary TKA, is to reduce pain and improve function. Revision TKA has been shown to be successful in improving patient outcomes in a cost-effective manner, however, in comparison to primary TKA, it is more expensive, has a higher complication rate, and has lower quality of life outcome scores [7-9].

Revision TKA is technically demanding and is potentially complicated by multiple factors not present during primary TKA including the need to remove components, sepsis, scarring and arthrofibrosis, ligamentous insufficiency or compromise, bone defects, metallic and polyethylene wear debris with variable levels of osteolysis, and deformity. Modern revision TKA systems offer a high degree of modularity, offset, metallic augmentation, stem lengths and fixation methods, and degree of constraint (Table 1). The revision knee system evaluated in this study addresses these needs with a comprehensive interchangeability between femoral and tibial sizes, 360 degrees of femoral and tibial offset options to allow for component position optimization for the best load transfer, a high degree of varus/valgus constraint without the use of a hinged prosthesis, and a simplified trial first approach to revision to provide for a more efficient surgery. The early results of this retrospective database review are promising, with improvement in pain and functional scores with only one failure for aseptic means at early follow-up. Continued monitoring of this cohort is paramount to analyze midterm and long-term results.

Conclusion

The early results of this modular TKA revision system are promising for use in complex TKA, with only one aseptic failure observed. There has been substantial improvements in ROM and function in this cohort.

Table 1. Results of Constrained Total Knee Arthroplasty

Study	Number of Knees	Device	Description	Follow-up (years)	Survival
Donaldson III et al., CORR 1988 [10]	31	TC III	17 Primary, 14 Revision	3.8	80.6%
Chotivichit et al., J Arth 1991 [11]	27	TC III	9 Primary, 18 Revision	4.3	100%
Hohl et al., CORR 1991 [12]	61	TC III	Complex Primary & Revision	6.1	95.1%
Kavolus et al., J Arth 1991 [13]	16	TC III	5 Primary, 16 Revision	4.5	100%
Rand, J Arth 1991 [14]	21	TC III	Revision, severely damaged knees	4	85.7%
Rosenberg et al., CORR 1991 [15]	36	TC III	Revision	3.8	97.2%
Stern et al., CORR 1991 [16]	8	NA	Primary, valgus $\geq 10^{\circ}$	4.5	100%
Haas et al., JBJS Am 1995 [17]	19	ІВ ССК	Revision, uncemented stems	3.6	89%
Vince & Long, CORR 1995 [18]	13	ІВ ССК	Revision	6	75%
Lachiewicz & Falatyn, J Arth 1996 [19]	46	TC III & Constrained Condylar	25 Primary, 21 Revision	46	95.7%
Lombardi et al., Seminars in Arthroplasty 1996 [20]	66	Maxim PSC	Revision	2	92.4%
Font-Rodriguez et al., CORR 1997 [21]	64	Constrained Condylar	Primary, cemented	7	98.1%
Peters et al., J Arth 1997 [22]	43	TC III, Constrained Condylar, Maxim PSC	Revision, cemented	5.2	97.7%
Hartford et al., J Arth 1998 [23]	33	TC III	17 Primary, 16 Revision, 8 Reimplantation	5	91%
Easley et al., CORR 2000 [24]	44	IB CCK	Primary for valgus deformity	7.8	100%
Nazarian et al., CORR 2002 [25]	207	IB CCK	Revision, 55 no stems	4.7	92.8%
Sheng et al., JBJS Br 2005 [26]	16	TC III	Revision, inflammatory	6.2	81.3%
Anderson et al., CORR 2006 [27]	55	Optetrak CCK	Primary without stems for valgus deformity	3.6	100%
Berend et al., CORR 2006 [28]	5	Maxim PSC	Primary, flexion contracture >20°	3.1	100%
Lachiewicz & Soileau, J Arth 2006 [29]	54	IB CCK & TC III	Primary	10	96%
Lombardi & Berend, Orthopedics 2006 [30]	421	Maxim PSC	47 Primary, 284 Revision, 90 Reimplantation	5	88.6%
Sheng et al., Acta Orth 2006 [31]	71	TC III	Revision	5.9	91.5%
Anderson et al., J Knee Surg 2007 [32]	248	Optetrak CCK	Primary without stems	3.9	97.5%
Lombardi et al., JBJS Am 2007 [33]	61	Maxim PSC	Complex Primary	5.6	88.5%
Kim & Kim, JBJS 2009 [34]	114	LCCK	Revision	7.2	96%
Peters et al., J Arth 2009 [35]	184	Maxim PSC & Vanguard PSC	Revision, cemented with cementless stems	4.1	91.8%
Hwang et al., COS 2010 [36]	25	LCCK	Revision, cemented	2.5	100%
Lachiewicz & Soileau, J Arth 2011 [37]	27	LCCK	Primary	5.4	96%
Lee et al., KSSTA 2013 [38]	79	LCCK	Revision	5.3	89.9%
Lee et al., J Arth 2013 [39]	27	LCCK with PS insert	Revision	7.0	100%
Pang et al., KSSTA 2013 [40]	50	LCCK & PFC Sigma TC III	Primary, type II valgus	8.3	96%

TCIII = Total Condylar III (Depuy, a Johnson & Johnson company, Warsaw, IN); NA = not available; IB CCK = Insall-Burstein Constrained Condylar Knee (Zimmer, Warsaw, IN); Constrained Condylar (Zimmer); Maxim PSC = Maxim Posterior Stabilized Constrained (Biomet, Warsaw, IN); Optetrak CCK = Optetrak Constrained Condylar Knee (Exactech, Gainesville, FL); Vanguard PSC = Vanguard Posterior Stabilized Constrained (Biomet); LCCK = Legacy Constrained Condylar Knee (Zimmer); PFC Sigma = Press-Fit Condylar Sigma (DePuy).

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