A Case Report of Revision Total Knee Arthroplasty After 17 Years: All Grown Up, What Happens When Implants Mature?

Colin K. Cantrell 1 , Harshadkumar A. Patel 2 , Wesley R. Stroud 3 , Nicholas Dahlgren 4 , Eva Lehtonen 5 , Morad Qarmali 6 , Kelly C. Stéfani 7 , Ashish Shah 2 , Sameer M. Naranje 2

 UAB School of Medicine, University of Alabama at Birmingham, Birmingham, USA 2. Orthopaedic Surgery, University of Alabama at Birmingham, Birmingham, USA 3. Orthopaedic Surgery, University of Alabama at Birmingham, Mountain Brook, USA 4. School of Medicine, University of Alabama at Birmingham, Birmingham, USA 5. Miller School of Medicine, University of Miami, Miami, USA 6. Pathology, University of Alabama at Birmingham, Birmingham, USA 7. Orthopaedic Surgery, Hospital Do Servidor Público Estadual, Sao Paulo, BRA

Corresponding author: Sameer M. Naranje, sameernaranje@gmail.com

Abstract

The number of total knee arthroplasties (TKAs) being performed annually is steadily rising. Recommendations for clinical follow-up guidelines following these arthroplasties is controversial, with no strict guidelines for long-term follow up. Although a few case series exist which identify a minority of patients who require revision TKA for aseptic loosening or pain more than 15 years after index surgery, no published studies have yet described these patients or the pathology present at the time of surgery in detail. We present the case of a patient who underwent revision TKA for pain and instability that developed 17 years after index surgery. Postoperative pathology revealed foreign body giant cell reaction of the tissue surrounding the previous implant. This case of revision after more than 17 years attempts to improve our understanding of long-term reactions to implants and highlights the necessity of long-term follow up in patients with TKA. It is one of the longest follow-ups of TKA reporting long-term anatomic changes at the bone cement interphase and around the implant.

Categories: Orthopedics, Other

Keywords: revision tka, loosening, follow-up, case report, total knee arthroplasty

Introduction

For more than a decade, the number of total knee arthroplasties (TKAs), primary and revision, has been steadily increasing [1]. With approximately 8% of them being revision TKAs, the documented reasons and timing for these revisions vary greatly [1-2]. Clinical follow up after TKA is controversial, and with the American Association of Hip and Knee Surgeon's (AAHKS) current recommendation of follow up annually or biennially with no definitive end date, adherence to these guidelines is questionable [3-4]. The longest documented follow up ranges up to 25 years, although complications may occur past this point [5]. One such complication, as presented in this case, is foreign body giant cell reaction [6].

Case Presentation

An 83-year-old female with bilateral primary TKA performed 17 years prior presented to the clinic. The patient was referred with worsening left knee pain, reported gait instability, and swelling for three months duration. Until this point, she had been completely asymptomatic. She was initially seen and treated by an orthopaedic surgeon from an outside facility with physical therapy, followed by a left knee arthrocentesis to rule out infection. The aspirate demonstrated proteinaceous fluid with few benign inflammatory and epithelial cells and cultures were found to be negative. Due to the increasing pain, gait instability, and discomfort, coupled with lack of relief by the current measures, she was referred to the orthopaedic surgery clinic at our institution for further evaluation.

At her initial visit, the patient reported steadily increasing, sharp pain localized to her left knee joint with associated swelling that worsened with ambulation and prolonged standing and lacked improvement with conservative management. Her day-to-day activities were becoming restricted secondary to the pain and she reported occasional falls due to the perceived instability of her knee joint. Physical exam revealed a mild antalgic gait and tenderness over her proximal tibia. An in-house X-ray was notable for an increase in size and number of osteochondral bodies in the left suprapatellar recess with a left joint effusion and "lysis and subsidence of the tibial component and decreased thickness, suggestive of loosening and wear" (Figures *1-4*). The patient then underwent a bilateral knee bone scan which confirmed the tentative diagnosis of implant loosening with polyethylene wear and instability. The patient was counseled on her treatment options, including surgical and non-surgical management, and elected to undergo revision surgery of her left knee arthroplasty.

Received 06/04/2018 Review began 06/06/2018 Review ended 06/08/2018 Published 06/13/2018

© Copyright 2018

Cantrell et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 3.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

How to cite this article

Cantrell C K, Patel H A, Stroud W R, et al. (June 13, 2018) A Case Report of Revision Total Knee Arthroplasty After 17 Years: All Grown Up, What Happens When Implants Mature?. Cureus 10(6): e2797. DOI 10.7759/cureus.2797



FIGURE 1: Standing radiograph of the case at initial presentation showing lysis and subsidence of left tibial component



FIGURE 2: Radiograph of the case at initial presentation showing osteochondral bodies in left suprapatellar recess



FIGURE 3: Anteroposterior radiograph of left knee at presentation showing lysis and subsidence of tibial component



FIGURE 4: Lateral radiograph of left knee at presentation showing large osteochondral bodies in suprapatellar recess

A classical anterior approach to the knee was made through the patient's previous scar. A medial parapatellar arthrotomy was performed, after which the knee joint was exposed, revealing extensive osteophytes around the patella (Figure 5). The osteophytes were removed and a medial release was performed, allowing for removal of the previous components (Figure 6). The femoral component was found to have bone ingrowth, which had encased the patella (Figure 7). Visualization of the bone-cement interface intra-operatively proved difficult. Upon gross visual inspection, it appeared that local long-term reaction at bone-cement interface had engulfed the cement and resulted in direct ingrowth of bone to implant. Bone-implant interface tissue was taken for histology examination. Microscopically, the sections examined showed papillary synovial proliferation which is consistent with the patient's history of long-standing osteoarthritis (Figure 8). Multiple foreign body giant cells, which are formed by fused macrophages, are seen in response to polarizable foreign material (Figure 9). Orthopaedic implants can cause chronic inflammation and giant cell foreign body response as seen in this case (Figure 10).

Cureus

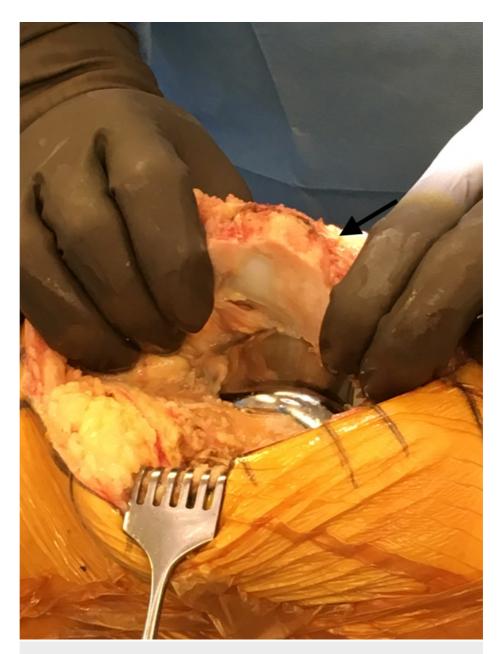


FIGURE 5: Intraoperative image of patella upon exposure showing extensive osteophyte development



FIGURE 6: Intraoperative image showing removal of previous total knee arthroplasty (TKA) components

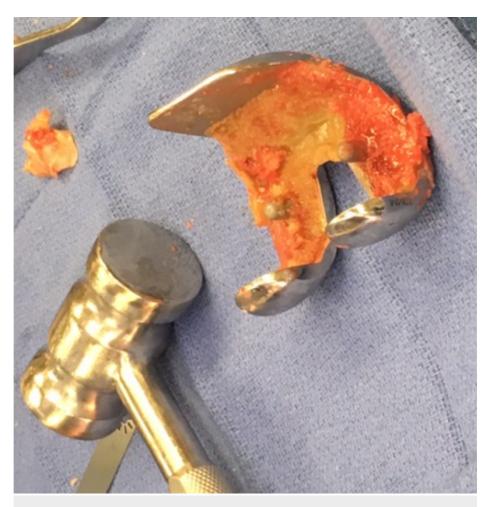


FIGURE 7: Intraoperative image showing bone ingrowth on femoral component of the previous implant

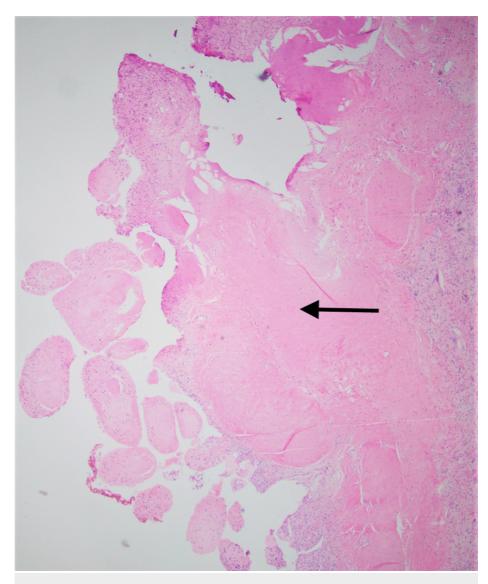


FIGURE 8: Low power photomicrograph showing papillary synovial proliferation; hematoxylin and eosin (H&E) stain

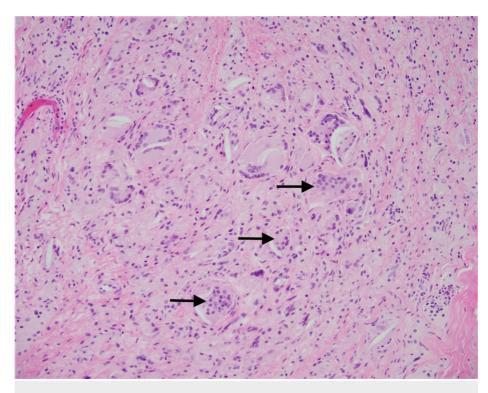


FIGURE 9: High power photomicrograph showing multiple foreign body giant cells in response to polarizable foreign material; hematoxylin and eosin (H&E) stain

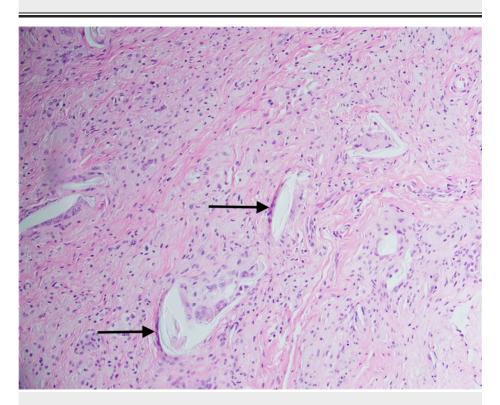


FIGURE 10: A photomicrograph showing multiple polarizable foreign material being engulfed by giant cells; hematoxylin and eosin (H&E) stain

After component removal, joint preparation was done in standard fashion by membranous tissue removal

and minimal freshening of previous bone cuts. Then revision of knee components was performed as per standard technique.

Postoperative X-rays confirmed excellent placement of a left knee arthroplasty (Figures *11-12*). The patient experienced no postoperative complications and was discharged from the hospital on postoperative day 2. She then followed up in the clinic two weeks postoperatively. The patient stated her pain was well controlled and had been working well with physical therapy. X-rays performed at this time reported that the left knee arthroplasty was in expected position with no evidence of hardware failure or loosening (Figures *13-14*). She reported that she was pleased with her new prosthesis.



FIGURE 11: Postoperative anteroposterior radiograph confirming proper placement of left revision total knee arthroplasty (TKA) components



FIGURE 12: Postoperative lateral radiograph confirming proper placement of left revision total knee arthroplasty (TKA) components





FIGURE 13: Postoperative radiograph at two-week follow up

FIGURE 14: Postoperative radiograph at two-week follow up

Discussion

Several indications for revision TKA have been defined in literature including mechanical wear, aseptic loosening, infection, instability, osteolysis, pain, stiffness, malalignments, patellar complication, and periprosthetic fractures [7-8]. The most common indication for TKA revision within two years of surgery is infection [9]. However, implant instability is the leading overall cause for the need of revision surgery possibly due to the increasing probability of implant instability in the aging implant, as well as the decreasing age of patients receiving primary TKA [10-12]. From 1991-2011, 30% of primary TKAs were performed in patients aged less than 65 years [12]. This younger, more active patient population is at an increased risk for wear-induced osteolysis [13]. These patients also have an increased period for immune system mediated reactions to occur, potentially loosening the prosthesis [14].

Foreign body giant cell (FBGC) reaction is not commonly observed or described in knee arthroplasty. Giant cells are formed by the fusion of cells from the monocyte/macrophage lineage [6,15]. Cells of this lineage are among the first cells to come in contact with the implanted biomaterials [16]. These FBGCs are most commonly observed at the interface of tissue and implanted material, such as knee arthroplasty prosthesis in this case [6]. The purpose of these giant cells mirrors that of osteoclasts: degradation and resorption of the underlying substrate. The main difference between these two cell types is that osteoclasts adhere to the bone while FBGCs adhere primarily to synthetic surfaces [6].

Sheikh et al. examined osseointegration, the term given to the foreign body responses to implant surfaces [15]. This osseointegration, a direct result of the foreign body reaction, may allow for bone to grow over the implant surface given the right conditions. This process, while commonly identified in dental implants, has been rarely noted in the context of knee arthroplasty. Our histology reports suggest that the same kind of reaction also occurred in this patient, leading to revision knee arthroplasty. Harkel et al. demonstrated that multinucleated giant cells (MNGC) are unable to resorb bone with the same capabilities of osteoclasts [17]. However, the ability of MNGC's to reabsorb hydroxyapatite, as well as to enhance osteoclast ability by more than 40-fold, was demonstrated.

The exact mechanism of this reaction of FBGCs and osteoclasts on bone, prostheses, and hydroxyapatite cement is incompletely understood [6]. Cytokine profiles in macrophages and FBGC's in arthroplasty tissue are similar to those of mature osteoclasts [14]. However, what determines the differences in the course and progression of these reactions are unknown [18]. Although the mechanisms behind this reaction are incompletely understood, periodic radiological follow up may identify osteogenesis and osteolysis occurring around the arthroplasty. Further studies would contribute to a greater understanding of this pathological process.

AAHKS recommendation for clinic radiological follow up is annually for the first five years, biennial for years 6-10, and then either annual or biennial for the years thereafter [3]. However, the recommendations varied according to physician group sizes, patient population sizes, and the total number of surgeries performed, and adherence to guidelines in clinical practice is far from universal. De Pablo et al. reported that less than half of patients who received a total hip arthroplasty had consistent radiographic follow-up [19].

Joshi et al. examined a large number of joint arthroplasty patients who were lost to follow up and perceived that these patients had no issues with their new joint [20]. This study implied that patients who experience issues and require revision present to their surgeon with symptoms leading towards that diagnosis. However, in some patients, by the time symptoms arise, there may be an extensive destruction of periprosthetic bone [12]. Clinical symptoms may be preceded by radiologic signs of implant failure [12]. Early detection of implant failure by routine follow up may prevent the need for complex revision surgery, which is more costly and laborious process to the patient.

Conclusions

FBGC and osteoclastic activity contributing to loosening is often a cause for component loosening in joint replacement surgeries. The fact that this patient remained clinically asymptomatic for over 15 years, until the aseptic osteolysis and component loosening became severe, highlights the importance of long-term follow up. With an increasingly younger patient age at the time of joint arthroplasty, long-term follow up to assess for signs and symptoms of implant failure is crucial for early diagnosis and to avoid more complex revision surgeries.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. N/A issued approval N/A. This study was exempt from IRB approval. . **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Kurtz S, Ong K, Lau E, Mowat F, Halpern M: Projections of primary and revision hip and knee arthroplasty in the United States from 2005 to 2030. J Bone Joint Surg Am. 2007, 89:780-785.
- Niinimaki TT: The reasons for knee arthroplasty revisions are incomparable in the different arthroplasty registries. Knee. 2015, 22:142-144. 10.1016/j.knee.2014.12.007
- Teeny SM, York SC, Mesko JW, Rea RE: Long-term follow-up care recommendations after total hip and knee arthroplasty: results of the American Association of Hip and Knee Surgeons' member survey. J Arthroplasty. 2003, 18:954-962.
- 4. Lovelock TM, Broughton NS: Follow-up after arthroplasty of the hip and knee: are we over-servicing or under-caring?. Bone Joint J. 2018, 100:6-10. 10.1302/0301-620X.100B1.BJJ-2017-0779.R1
- Patil S, McCauley JC, Pulido P, Colwell CW, Jr: How do knee implants perform past the second decade? Nineteen- to 25-year followup of the Press-fit Condylar design TKA. Clin Orthop Relat Res. 2015, 473:135-140. 10.1007/s11999-014-3792-6
- Brodbeck WG, Anderson JM: Giant cell formation and function. Curr Opin Hematol. 2009, 16:53-57. 10.1097/MOH.0b013e32831ac52e
- Tay KS, Lo NN, Yeo SJ, Chia SL, Tay DK, Chin PL: Revision total knee arthroplasty: causes and outcomes. Ann Acad Med Singapore. 2013, 42:178-183.

- Sharkey PF, Hozack WJ, Rothman RH, Shastri S, Jacoby SM: Insall Award paper. Why are total knee arthroplasties failing today?. Clin Orthop Relat Res. 2002, 7-13.
- 9. Meding JB, Ritter MA, Davis KE, Farris A: Meeting increased demand for total knee replacement and followup: determining optimal follow-up. Bone Joint J. 2013, 95:1484-1489. 10.1302/0301-620X.95B11.32467
- Singh J, Politis A, Loucks L, Hedden DR, Bohm ER: Trends in revision hip and knee arthroplasty observations after implementation of a regional joint replacement registry. Can J Surg. 2016, 59:304-310.
- 11. Khan M, Osman K, Green G, Haddad FS: The epidemiology of failure in total knee arthroplasty: avoiding your next revision. Bone Joint J. 2016, 98:105-112. 10.1302/0301-620X.98B1.36293
- Bayliss LE, Culliford D, Monk AP, et al.: The effect of patient age at intervention on risk of implant revision after total replacement of the hip or knee: a population-based cohort study. Lancet. 2017, 389:1424-1430. 10.1016/S0140-6736(17)30059-4
- Clohisy JC, Kamath GV, Byrd GD, Steger-May K, Wright RW: Patient compliance with clinical follow-up after total joint arthroplasty. J Bone Joint Surg Am. 2008, 90:1848-1854. 10.2106/JBJS.G.00856
- 14. Neale SD, Athanasou NA: Cytokine receptor profile of arthroplasty macrophages, foreign body giant cells and mature osteoclasts. Acta Orthop Scand. 2009, 70:452-458. 10.3109/17453679909000980
- 15. Sheikh Z, Brooks PJ, Barzilay O, Fine N, Glogauer M: Macrophages, foreign body giant cells and their response to implantable biomaterials. Materials. 2015, 8:5671-5701. 10.3390/ma8095269
- 16. Miron RJ, Zohdi H, Fujioka-Kobayashi M, Bosshardt DD: Giant cells around bone biomaterials: osteoclasts or multi-nucleated giant cells?. Acta Biomater. 2016, 46:15-28. 10.1016/j.actbio.2016.09.029
- ten Harkel B, Schoenmaker T, Picavet DI, Davison NL, de Vries TJ, Everts V: The foreign body giant cell cannot resorb bone, but dissolves hydroxyapatite like osteoclasts. PLoS One. 2015, 10:0139564. 10.1371/journal.pone.0139564
- Ahmed GJ, Tatsukawa E, Morishita K, et al.: Regulation and biological significance of formation of osteoclasts and foreign body giant cells in an extraskeletal implantation model. Acta Histochem Cytochem. 2016, 49:97-107. 10.1267/ahc.16007
- 19. de Pablo P, Losina E, Mahomed N, et al.: Extent of followup care after elective total hip replacement. J Rheumatol. 2006, 33:1159-1166.
- 20. Joshi AB, Gill GS, Smith PL: Outcome in patients lost to follow-up. J Arthroplasty. 2003, 18:149-153. 10.1054/arth.2003.50061