

# Polyethylene Insert Wear Leading to Early Aseptic Bilateral Total Knee Arthroplasty Failure: Case Report

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## Learning Point of the Article:

Challenges faced post-TKA implant failure and how to address them.

## Abstract

**Introduction:** Component failure is one of the most dreaded complications of any total knee arthroplasty (TKA) surgery. To tackle this, implant technology continues to evolve with the goal of increasing implant survivorship post-TKA surgery. It is estimated that about 20% of TKA revisions are due to mechanical aseptic loosening. Polyethylene insert wear has become an increasingly common finding at the time of revision surgeries. With this case report, we try to highlight the challenges faced during bilateral revision TKA and the use of both femoral and tibial augments to compensate for the bony osteolysis.

**Case Report:** A 69-year-old patient with unusual presentation of bilateral TKA (Exactech, Hiflex) failure 6 years post-surgery due to polyethylene wear. Three months following surgery, the patient had no complications and had satisfactory radiological, pain, and functional outcomes.

**Conclusion:** Although aseptic TKA failure due to polyethylene wear is an uncommon finding. Our case of the 69-year-old male with bilateral arthroplasty failure highlights the significance of the durability of polyethylene along with its wear and fracture resistance. It demonstrates that timely intervention with revision arthroplasty drastically improves pain, mobility, joint stability, and overall quality of life in these patients.

**Keywords:** Polyethylene wear, revision arthroplasty, total knee arthroplasty failure, augments, case report.

## Introduction

Total knee arthroplasty (TKA) has become one of the most frequently performed orthopedic procedures worldwide. It is estimated that more than 750,000 TKA cases were performed in 2014 [1]. In a study by Inacio et al. [2] in 2017, TKA number is projected to increase annually by 143% until 2050, translating to a projected volume of 1.5 million TKAs in the United States alone. The total number of knee revision surgeries is expected to grow by 601% by 2030 [3]. TKA component failure is one of the

most dreaded complications of a replacement surgery. Therefore, implant technology continues to evolve to increase implant survivorship post-TKA surgery. A recent study demonstrated 96.1% and 89.7% survivorship at 10 and 20 years of TKA, respectively [4]. It is estimated that about 20% of TKA revisions are due to mechanical aseptic loosening [5]. Although many etiologies and factors can lead to mechanical loosening, one specific cause that has recently drawn more attention is the debonding of the tibial, femoral, or both components at the cement-implant interface [6]. One of the causes of this

## Author's Photo Gallery



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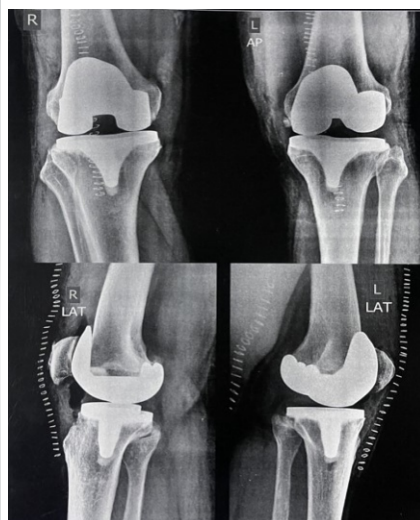
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**Figure 1:** Pre-operative B/L Knee weight-bearing AP/Lat X-ray.



**Figure 2:** Post-operative day 1 B/L Knee AP/Lat X-ray.



**Figure 3:** 6 years post-operative B/L Knee AP/Lat X-ray showing implant loosening.

debonding is the wear of the polyethylene insert, which increases the stress on the cement-implant interface.

When implant failure occurs, it can lead to persistent knee pain, recurrent effusion, and subsequent component migration, which may require partial or complete revision surgery [7]. This type of mechanical debonding may be challenging to diagnose, as radiographs often do not show clear evidence of aseptic loosening, and migration can be a late finding [8]. According to Rao et al., wear occurring at the interface between the polyethylene insert and the modular tibial baseplate has become an increasingly common finding at the time of TKA revision [9]. The longevity of the polyethylene tibial inserts also depends on the sterilization method. There have been

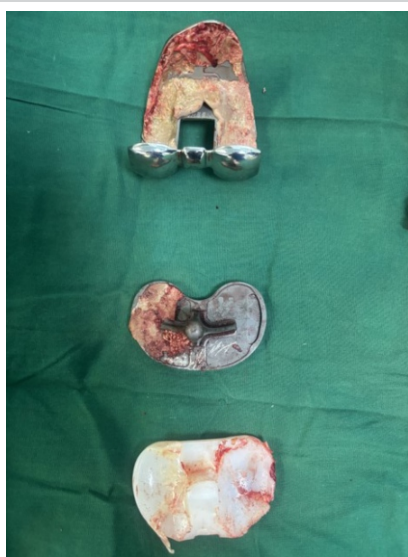
increasing concerns regarding bearings sterilized with radiation in an inert gas may lead to oxidization

in vivo and develop fatigue wear. This wear can be attributed to free radicals generated during sterilization with radiation [10]. At the same time, non-irradiated bearings may undergo more significant losses in thickness from routine burnishing since they lack the cross-linking that accompanies sterilization with radiation [10].

This case discusses an unusual presentation of bilateral TKA (Exactech, Hiflex) failure 6 years post-surgery due to polyethylene wear leading to implant loosening. With this case report, we highlight the challenges faced during the bilateral revision of TKA and the use of both femoral and tibial augments to compensate for the bony defect.

### Case Report

A 69-year-old, right-side dominant patient of Kellgren and Lawrence Pai et al. [11] grade 4 osteoarthritis (Fig. 1) underwent (Exactech, Hiflex) bilateral posterior stabilized primary TKA (Fig. 2) in a private hospital. The patient did not have any complaints and had been functioning well following his arthroplasty. 5 years after primary surgery, the patient presented with a history of bilateral knee pain (Right > Left) and joint instability, gradually increasing over the past 8 months. The patient also had difficulty walking with a painful limp. There was no history of any fall or trauma. The patient is a non-diabetic and normotensive



**Figure 4:** Intraoperative images of explanted total knee replacement components.



**Figure 5:** Polyethylene wear on the medial side of the insert.





**Figure 6:** B/L Revision total knee arthroplasty post-operative X-ray.

individual. Surgical error as a possible causative factor was excluded because the patient had been functioning well after surgery.

On examination of both knees, effusion was noted along with joint line tenderness. Range of motion was 0–80° on the right knee and 0–110° on the left knee. Mediolateral instability was noted during the examination. The neurological and systemic examination revealed normal findings. The

overall Visual Analog Scale (VAS) score [12] 3 months post-primary surgery was 1; this score increased significantly 9 to 6 years post-primary surgery. Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) score [13] for the right knee three months post-primary surgery was 6 and 2 for the left knee. Six years post-primary surgery, the WOMAC score increased to 73 for the right knee and 45 for the left knee. For the right knee, three months post-primary surgery, the Forgotten Joint score (FJ. Score) [14] was 89.6 and 91.7 for the left knee. At the same time, FJ. score 6 years post-primary surgery was 14.6 for the right knee and 46.7 for the left knee.

### Radiological findings

6 years post-operative X-ray (Fig. 3) findings, including knee

standing A/P and lateral radiograph, revealed the femoral component loosening more on the right side. The A/P view of both knees revealed reduced joint space and polyethylene wear on the medial side. The lateral view indicated resorption of the posterior femoral condyles and debonding at the cement-bone interface more on the right than the left side.

### Laboratory findings

The fluid culture from both knees showed no signs of infection. Gram and ZN stains were negative but revealed increased levels of neutrophils. Other basic investigations were normal except C-reactive protein, which was weakly positive.

### Surgical procedure

#### Stage one

Right knee was operated on first, given being more symptomatically involved. Anterior midline skin incision was taken over the previous surgical scar. For arthrotomy, a medial para-patellar approach was used for the right knee. Synovitis was observed over the anterior femoral condylar surface. Both femoral and tibial implant loosening with polyethylene wear was noted. The polyethylene insert was detached from the tibial base plate using an osteotome. A loose femoral component, utterly detached from the femoral condylar surface, was noted. With the help of osteotome, an interface was created between the tibial bone and cement (Fig. 4). Diffuse osteolysis was noted over the medial femoral and tibial condyles. A synovectomy was done, and the fibrosis was released. The tibia and femoral preparation was done, and bony cuts were made. Flexion and Extension gap balancing using trial implants showed tight extension gaps. Hence, the bony cuts were revised, and the final gap balancing was done using trial implants.

Two 5 mm distal femoral augments were added to the trial femoral component to overcome the bony defect. Flexion and Extension gap balancing were re-done using trial implants. The

### AORI CLASSIFICATION TKA BONE LOSS

TYPE	DESCRIPTION	FIXATION
1	intact metaphyseal bone, contained defect	standard TKA
2	cortical bone loss in metaphysis (uncontained)	
2A	one femoral/tibial condyle	< 5 mm defect - cement only
2B	both femoral/tibial condyles	5 - 10 mm defect - cement + rebar
		> 10 mm defect - augments
3	significantly deficient metaphysis	bypass metaphysis for stability: cone/sleeve, or oncology prosthesis

**Figure 7:** Anderson orthopedic research institute classification for total knee arthroplasty bone loss.

final right knee implantation was done using a Maxx progressive constraint knee (PCK) freedom implant with femoral component size C with an extension stem of  $15 \times 75$  mm. Two distal femoral augments and two posterior femoral augments of 5 mm each were used. The offset junction was set at 4mm. A tibial component of size 4 with an extension stem of  $12 \times 40$  mm was selected and set at an offset junction of 4 mm. The bony defect on the tibial surface was built with bone grafts and cement. It must be noted that both the femoral and tibial canals were filled with cement before implantation. The final polyethylene insert of 17 mm PCK was implanted. All of the extravasated cement was meticulously removed before the hardening of the cement. A wound wash was given, and closure was done in layers over a suction drain after achieving adequate hemostasis.

## Stage 2

The left knee was operated on 2 days following the right knee revision. Anterior midline skin incision was taken over the previously operated scar. Following medial para-patellar arthrotomy, the knee joint was exposed. The femoral component was not as loose as that of the right side. Synovitis was noted over the anterior femoral surface. The polyethylene insert and femoral and tibial components were removed. Polyethylene wear was also noted (Fig. 5). Osteolysis was observed over the medial femoral and tibial condyles. The defect over the medial tibial condyle was more significant than that of the opposite knee. Synovectomy was done along with fibrosis release. After taking the bony cuts, the tibia and femoral preparation were done.

Flexion and Extension gap balancing were done using trial implants and augments. Left revision TKA was done using Maxx PCK freedom implant. The final inserted femoral component was size C with a  $12 \times 40$  mm extension stem. Two posterior femoral augments of 5 mm each were used, and the offset junction was set at 4 mm. The tibial component of size 4 was selected and fixed with an extension stem of  $13.5 \times 75$  mm set at an offset junction of 4 mm. A tibial augment of 5 mm was attached to build the bony defect over the medial tibial condyle. The polyethylene insert of 14 mm PCK was inserted. The cementing technique was similar to that of the right knee. Wound wash was given, and a suction drain was inserted. Wound closure was done in layers after achieving hemostasis.

Intraoperative cultures taken from both knees were negative.

## Postoperative result

The patient had a successful and uncomplicated postoperative rehabilitation and was released on postoperative day 5. All

laboratory investigations were within normal limits at the time of discharge. Oral analgesics and physiotherapy facilitated a progressive recovery. For the initial 3 weeks, the patient used a walker for ambulation. Three weeks following surgery, the patient ambulated without any assistance. The patient resumed his occupation after 2 months of rehabilitation. Three months following surgery, the overall VAS score for pain was 0; the WOMAC score improved to 7 on the right knee and 4 on the left knee.

Furthermore, the F.J. score was 82.4 for the right knee and 86.8 for the left knee. The postoperative X-ray of the bilateral knees acquired in standing A/P and lateral position revealed integrated components with a well-balanced joint line (Fig 6).

## Discussion

With final-stage knee osteoarthritis, TKA is the treatment of choice [15]. Patient selection and pre-operative diagnosis affect the outcome of aseptic TKA revision surgery [16]. Patients with aseptic failure can perform quite well if treated with proper surgical technique and appropriate implant selection [17]. According to Friedman et al., the functional improvement in stability, motion, and pain is significantly less after revision surgery for aseptic failure than after primary TKA [18]. Patients with aseptic failure but without knee stiffness have been noted to have a better outcome [16].

Meanwhile, patients who undergo revision TKA for stiffness instead of loosening or instability have inferior outcomes [19]. It is to be noted that aseptic revision TKA achieved a significantly better Knee Society Score and range of motion. In contrast, pain and functional scores were similar in septic and aseptic revision surgery [16].

This case is a peculiar presentation of polyethylene wear, leading to bilateral TKA failure. Orthopaedic practice has evolved from cement-on-cement articulating spacers to the routine use of metal-on-polyethylene spacers [20]. Therefore, the durability of the polyethylene, along with its wear and fracture resistance, are important factors for a successful arthroplasty outcome. For this patient, the revision of bilateral TKA with subsequent debridement and polyethylene component evaluation gave insight into the reasons for the catastrophic failure of the arthroplasty surgery. The bone loss due to osteolysis on the right and left knee was of type 2B as per the Anderson Orthopaedic Research Institute classification of TKA bone loss (Fig. 7) [21]. According to Rand in the research done at Mayo Clinic, Rochester, the knee scores drastically improved after the use of metal wedge augmentation for tibial bony defects [22]. Therefore, these osteolytic defects on tibial and femoral condyles in both right and left knees were

successfully overcome using bone graft, metal augments, and cement mantle.

arthroplasty drastically improves pain, mobility, joint stability, and overall quality of life in these patients.

### Conclusion

Although aseptic TKA failure due to polyethylene wear is an uncommon finding, our case of the 69-year-old male with bilateral arthroplasty failure highlights the significance of polyethylene's durability, along with its wear and fracture resistance. It demonstrates that timely intervention with revision

### Clinical Message

Research into the development of durable, wear-resistant, and fracture-proof polyethylene is essential for long-lasting TKA surgery. Tibial and Femoral augments must be used judiciously.

**Declaration of patient consent:** The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient has given the consent for his/ her images and other clinical information to be reported in the journal. The patient understands that his/ her names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

**Conflict of interest:** Nil    **Source of support:** None

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