Three-Year Clinical Results of a Cruciate-Retaining Type of the Knee Prosthesis with Anatomical Geometry Developed in Japan

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Abstract

**Background:** The FINE total knee was developed in Japan and clinical use began in 2001. It has unique design features, including an oblique 3° femorotibial joint line that reproduces anatomical geometry. Although 20 years have passed since the FINE knee was clinically used for the first time in Japan, a formal clinical evaluation including patient-reported and radiographic outcomes has not been undertaken.

**Methods:** A total of 175 consecutive primary cruciate-retaining (CR)-FINE total knee arthroplasties (TKAs) at our hospital between February 2015 and March 2017 were included in this study. Three years postoperatively, range of motion (ROM), Knee Society Score (KSS), Knee Injury and Osteoarthritis Outcome Score (KOOS) and Forgotten Joint Score (FJS) were recorded and compared with preoperative scores. Radiographic analyses including mechanical alignment, component alignment, and incidence of radiolucent lines also were undertaken based on the radiographs three years postoperatively.

**Results:** One-hundred twenty-two knees (70%) were available for 3-year follow-up data using KOOS, except for the sports subscale. Postoperative KOOS-symptom, -pain and -ADL were > 85 points, but KOOS-sports, -QOL and FJS were less satisfactory. ROM, KSS and all the subscales of KOOS were significantly improved compared with preoperative scores. Postoperative mean FJS was 66 and was significantly correlated with all the subscales of KOOS, but not with postoperative ROM. Radiolucent lines ≥1 mm wide were detected in five knees (4.1%). There were no major complications needing revision surgeries.

**Conclusions:** Patient-reported outcomes (PROs) for symptoms, pain and ADL after the CR-FINE TKA were generally improved, but those for sports, QOL and FJS were improved less. The incidence of radiolucent lines was rare but detected around the femoral components. Improvements of surgical technique or innovation of the implant design with mid- to long-term follow-up will be necessary to achieve better PROs from patients receiving the FINE knee.

**Background**

Although the outcomes of total knee arthroplasty (TKA) are generally acceptable, approximately 20% of patients have some complaints after TKA (1-3). The reasons for dissatisfaction after TKA remain poorly understood; however, failure of restoration of a physiological joint line has been suggested as a causative factor. In 2011, Bellemans introduced kinematically aligned (KA)-TKA as a surgical technique to realize a physiological joint line (4). With KA-TKA, the femoral and tibial components are implanted with mild varus limb alignment, relative to neutral alignment, in order to restore the physiological joint line to a pre-arthritic state. While good patient reported outcomes after KA-TKA have been reported (5-9), the longevity of polyethylene inserts and femoral and/or tibial components implanted with a varus alignment are a concern (10, 11).
The FINE total knee has been developed in Japan and used for approximately 20,000 TKAs of Japanese patients since 2001. It has unique design features, including an oblique 3° femorotibial joint line (Fig. 1). This feature enables the implant to reproduce anatomical geometry and allows the osteotomy to be performed perpendicular to the mechanical axis. The FINE total knee is also designed to guide internal movements of the tibia via medial pivotal rotation, thus permitting deeper flexion of the knee to better match the lifestyle needs of Japanese populations (12). The medial surface of the polyethylene insert has a convex curve which is designed to increase the rate of conformity to the femoral component, thereby enhancing internal rotation of the tibia. Conversely, the lateral surface has a flat surface that has been designed to allow femoral rollback, enhancing internal rotation of the tibia via medial pivotal motion (12). Hence, the design concepts of the FINE total knee facilitate KA-TKA via conventional osteotomy.

Although 20 years have passed since this implant was clinically used for the first time in Japan, a formal clinical evaluation including patient-reported and radiographic outcomes has not been undertaken. The aim of the present study was to evaluate the 3-year clinical results including patient-reported and radiographic outcomes in Japanese patients receiving a cruciate-retaining (CR) type of the FINE total knee.

**Methods**

**Patients**

A total of 175 consecutive primary TKAs using a CR type of the FINE total knee (Teijin-Nakashima Medical Co. Ltd., Okayama, Japan) in 157 patients at our hospital between February 2015 and March 2017 were included in this study. One patient (one TKA) died from TKA-unrelated causes. None received a revision. Of the remaining 174 TKAs (156 patients), 122 knees (111 patients, 70%) were available for 3-year follow-up data; the data for the Knee Injury and Osteoarthritis Outcome Score (KOOS) (13) except for a sports subscale and radiographs were available for all of those while data for KOOS-sports and the Forgotten Joint Score (FJS) (14) were available for 53 (30%) and 77 knees (44%), respectively (Fig. 2). There were nine men and 102 women, with a mean age of 72.3 years (29-89) at the time of surgery. The mean body mass index was 27.2 kg/m² (16.7-39.6). One hundred twelve TKAs were performed for osteoarthritis, seven for rheumatoid arthritis (RA), and three for osteonecrosis (Table 1).

This study was approved by the institutional review board at our institution. All activities were performed in accordance with the ethical standards set forth in the Declaration of Helsinki, and informed consent was obtained from all patients who participated in this study.

**Surgical procedures**

All TKAs were performed using the measured resection technique by anterior reference (15). Surgical approaches were chosen either mid-vastus or sub-vastus for varus knees, but the lateral parapatellar approach was used for valgus knees. A release of the deep fibers of the medial collateral ligament (MCL) was routinely performed for varus knees. Surgeries were performed using conventional instruments; that
is, distal femoral osteotomy was conducted perpendicular to the mechanical axis at a level 9-10 mm from
the farthest point of the medial condyle, and the posterior condyle was osteotomized parallel to the
surgical epicondylar axis (3°external to the posterior condylar line). A tibial osteotomy subsequently was
conducted perpendicular to the anatomical axis of the tibia. The cutting level was set 8-10 mm distal to
the convex of the lateral plateau. Following osteotomy, adjustments for soft tissue balancing were
performed before the implants were fixed to the bone with cement. Whether to replace the patellar
component or not depended on the surgeons’ decision; 80 knees received patellar replacement (66%,
Table 1). Patients were discharged three weeks after surgery when they were medically stable, with pain
controlled by oral analgesics and deemed by a physiotherapist to be mobilizing sufficiently to function
safely at home.

**Radiographic examinations**

Routine postoperative assessment included anteroposterior, lateral, and 60°skyline radiographs of the
knee, and full-length standing radiographs of both lower limbs. The anatomical axis (the angle subtended
by lines bisecting the medullary canals of the femur and the tibia) and the mechanical axis (the angle
subtended by lines connecting the center of the femoral head and the center of the femoral condyles, and
the center of the tibial plateau to the center of the talus) were measured from full-length standing
radiographs. The alignment of the components was assessed on AP radiographs of the knee using the
distal femoral valgus angle (DFVA, $\alpha$) and proximal tibial varus angle (PTVA, $\beta$), while the femoral flexion
angle (FFA, $\gamma$) and tibial slope (TS, $\sigma$) were measured on lateral radiographs. The mechanical alignment
was assessed by the hip-knee-ankle (HKA) angle based on the full-length standing radiographs with varus
alignment designated as positive (Fig. 3). These measurements were performed using the protocol of
Kilincoglu et al (16). Three independent observers (AN, MY, KY) examined radiographs for evidence of
anterior notching of the femur, component failure or subsidence, lucent lines, osteolysis, and heterotopic
ossification based on the standardized Knee Society radiological evaluation system (17).

**Clinical evaluation**

We used the Knee Society Score (KSS), which consists of a knee score (KSS-KS) and a function score
(KSS-FS), as an objective evaluation of knee function (18). In addition to the KSS, we used the Japanese
KOOS, an instrument of confirmed validity and reliability for patient-reported outcomes (PROs) based on
its cross-cultural adaptation (19). The KOOS consists of a total of 42 knee-related items, and each item
was scored from 0-4. Five KOOS subscales, including symptoms (KOOS-symptom), pain (KOOS-pain),
ADL (KOOS-ADL), sports/recreation (KOOS-sports), and quality of life (KOOS-QOL) were converted to 100
points (13). Furthermore, we investigated the FJS for 77 TKAs (72 patients). The range of motion (ROM)
was measured using a goniometer. Postoperative scores were compared with the preoperative scores.
Both intraoperative and postoperative complications were noted from the medical records.

**Statistical analysis**
The reliability of each radiographic measurement was assessed using intraclass correlation coefficients. All radiographic measurements in this study showed good reliability (all values > 0.8), and discrepancies were discussed until consensus was achieved. The paired t-test was used to compare 3-year postoperative with preoperative scores. Results were expressed as the mean (standard deviation, SD). Correlations among postoperative KSS, KOOS and FJS were analyzed by Pearson's correlation coefficients. Data analyses were performed using SPSS software, version 21 (SPSS Inc., Chicago, IL, USA) and p-values of < 0.05 were considered statistically significant.

Results

Clinical outcome scores

Mean postoperative KSS-KS and KSS-FS (SD) were 97 (4.9) and 76 (19), respectively (Table 2). Mean postoperative KOOS-symptom, -pain, -ADL, -sports and -QOL (SD) were 85 (14), 89 (13), 86 (14), 53 (29) and 70 (21), respectively (Table 3). Patients reported clinically and statistically significant improvements in the KSS-KS, KSS-FS, and all subscales of the KOOS (p < 0.001, Table 3). Mean postoperative FJS was 66 (minimum: 19, maximum: 100) (Table 3).

Mean postoperative flexion angle (SD) was 124 (13.3)°. The postoperative flexion angle increased significantly compared with the preoperative one (p < 0.001, Table 2). The mean improvement in ROM (SD) was 13 (19)° (Table 2). Seven knees had limited extension; six knees had an extension limitation between 10° and 20° and one knee had a limitation of 25°.

Correlations among postoperative KSS, KOOS and FJS

Correlations among postoperative KSS, KOOS and FJS were computed to investigate the relationships among these postoperative outcomes. FJS was correlated significantly with KSS-FS and all the subscales of KOOS, but not with postoperative flexion angle or ROM (Table 4).

Radiographic outcomes

Mean postoperative standing FTA, DFVA (α), PTVA (β), FFA (γ) and TS (σ) are shown in Table 5. Mean postoperative HKA angle (SD) was 2.3 (3.6)° with slightly varus alignment, and 74 knees (60.7%) were within ± 3° of the HKA angle.

Radiographic analyses three years postoperatively revealed no instances of osteolysis or subsidence. Radiolucent lines ≥1 mm wide were detected in five knees (4.1%), all of which occurred in zone 4 of the femoral components but were insignificant clinically. There was one knee with heterotopic ossification in the quadriceps (0.8%), but it was asymptomatic (Table 5).

Complications
One patient died due to a cause unrelated to TKA. One had a suspicious deep infection but joint fluid culture was negative for bacteria and the knee was not revised. There were eight partial tears of the popliteal tendon intraoperatively, all of which were sutured using nylon thread. One patient had an intraoperative avulsion of the superficial fibers of the MCL from its insertion to the tibia, which was reconstructed by suture and pull-out. One patient with RA had a medial subchondral fracture of the proximal tibia intraoperatively, which was fixed using a cancellous screw. There was one lateral supracondylar fracture intraoperatively in an RA patient and one anterior nothing without a periprosthetic fracture, but no additional surgeries were required. One patient with a severely deformed valgus knee (preoperative femorotibial angle: 152°) had transient peroneal nerve palsy postoperatively but had recovered fully by the 3-year follow-up (Table 6).

Discussion

The most characteristic point of the FINE knee is the design that reproduces the anatomical geometry, that is, a 3° obliquity built into the medial femorotibial surface in both coronal and axial planes. Here we showed that the postoperative mean DFVA was 98°, slightly more than that of the conventional prostheses. These features allow surgeons to perform KA-TKA without the femoral component implanted slightly in valgus and the tibial component implanted slightly in varus. Furthermore, the FINE knee adopts an ultra-high molecular weight polyethylene insert including vitamin E with antioxidant properties. As there is an increased risk of revision in KA-TKA using conventional prostheses (20, 21), these characteristic designs of the FINE knee are expected to show superior longevity to the conventional prostheses implanted in kinematic alignment.

The second characteristic point of the FINE knee is the polyethylene insert, which is dish-shaped medially and has a flat-surface laterally. This structure allows natural internal rotation of the tibia and roll-back of the lateral femoral condyle during flexion, leading to deep flexion. Here, the mean postoperative flexion angle (SD) and the improvement in ROM were 124 (13)° and 13°, respectively (Table 2). The correlation analyses among postoperative KSS, KOOS and FJS showed significant correlations between flexion angle and KSS-KS, and ROM and KSS-KS, while flexion angle and ROM did not show any significant correlations with KOOS subscales or FJS (Table 4). These results suggest that postoperative flexion angle does not have a significant impact on PROs if it reaches over 120°. Rather, stability during all ranges of motion may be a more important factor to achieve better PROs.

For all patients included in this study, we performed TKAs using a measured resection (MR) technique. van Lieshout et al. showed that the joint line was elevated after TKA using the MR technique (22). In addition, Luyckx et al., using cadaver knees, demonstrated that despite a well-balanced knee in full extension and at 90° of flexion, increased mid-flexion instability was evident in knees in which the joint line was raised (23). Because we cut femoral posterior condyles by anterior reference and used a CR-type for the patients who participated in this study, we might have implanted femoral components that were smaller than the anatomical anteroposterior length of the femoral condyles, and this may have caused
shortening of the posterior condylar offsets. This may also raise the joint line, which in turn, causes mid-flexion instability.

Recently, the FJS has been used to evaluate top-performing TKAs since it has a diminished ceiling effect (14). Parratte et al described a cohort of posterior-stabilized Zimmer LPS-Flex TKA (Zimmer Biomet, Warsaw, Indiana) with a mean FJS of 74 at a mean follow-up of 3.8 years (24), while Thomsen et al reported mean FJS for the Vanguard CR TKA (Zimmer Biomet) of between 44 and 59 from 1 to 4 years following mobile-bearing or CR TKAs (25). Moreover, excellent 5-year clinical results of the medial ball and socket TKAs were demonstrated by Katchky et al (26). According to their report, the mean postoperative flexion angle was 124°. It was surprising that postoperative KOOS-symptom, -pain and -ADL were > 90, with KOOS-sports being 71 and KOOS-QOL being 82. Furthermore, postoperative Oxford Knee Score and FJS were 44 (full score: 48) and 75, respectively. These results are better than the results obtained from patients receiving a CR-type of the FINE knee. Although a simple comparison between the two different prostheses should be avoided as the backgrounds of the patients were different, these excellent mid-term results of the medial ball and socket knee suggest the importance of the congruent medial articulation.

To obtain better PROs from the FINE knee, we are now performing TKAs using the pre-cut technique developed by Kaneyama et al (27), which allows the amount of cutting of the femoral posterior condyles to the extension gap to be adjusted. These improvements in surgical technique are expected to avoid shortening of the posterior condylar offset, leading to better ROM and stability in flexion after TKA (28-30). Although we could not show better PROs for the CR-FINE knee than other top-performing prostheses developed in the US or Western Europe, these improvements in the surgical technique will raise the PROs after the CR-FINE TKA.

Radiographic analyses for the CR-FINE TKA three years postoperatively demonstrated radiolucent lines more than 1 mm wide in five knees (4.1%) (Table 6). All radiolucent lines were detected in zone 4 on the femoral component side. The possible reason for the radiolucent line in this location may be the application of less or no bone cement in that zone to avoid spill-over after implantation of the femoral components. Although no clinical problems occurred over three years postoperatively, continuous attention should be paid to whether these radiolucent lines will spread or not. It should be noted that no subsidence of the tibial components occurred during the 3-year follow-up period. Tibial base plates of the FINE knee adopt an asymmetric design between medial and lateral sides, which provides good coverage on the cut surface of the tibia. These features probably contribute to the absence of subsidence of the tibial components.

There are some limitations to this study. First, the sample size was relatively small and the patients were recruited from a single institution. Second, 30% of the knees were not available for analysis of 3-year KOOS data. Moreover, only 30% of knees were available for a KOOS-sports subscale, and 44% were available for FJS. Nevertheless, our results show that the FINE knee is widely acceptable for Japanese
patients with knee deformities, considering the good PROs comparable to other top-performing knee prostheses.

Conclusions

We showed 3-year clinical results of a CR-type of the FINE total knee that was implanted through the MR technique. Postoperative KOOS-symptom, -pain and -ADL were > 85 points, but KOOS-sports, -QOL and FJS were less satisfactory. There were no major complications needing revision surgeries. Incidence of radiolucent lines was rare around the femoral components. There were no instances of osteolysis or subsidence. Improvements of surgical technique or innovation of the implant design with mid- to long-term follow-up will be necessary to achieve better PROs from patients receiving the FINE knee.

Abbreviations

CR: cruciate-retaining

TKA: total knee arthroplasty

KSS: Knee Society Score

KOOS: Knee injury Osteoarthritis Outcome Score

PROs: patient-reported outcomes

FJS: Forgotten Joint Score

KA: kinematic alignment

ROM: range of motion

FTA: femorotibial angle

HKA: hip-knee-ankle

DFVA: distal femoral valgus angle

PTVA: proximal tibial varus angle

FFA: femoral flexion angle

TS: tibial slope

Declarations

Ethics approval and consent to participate
Approval for the study was received from the Institutional Review Board at Toho University Sakura Medical Center. Written informed consent was obtained from all the patients before surgery. All activities were performed in accordance with the ethical standards set forth in the Declaration of Helsinki.

Consent for publication

Not applicable.

Availability of data and materials

All data generated or analyzed during the current study are included in this published article.

Competing interests

The authors declare that they have no competing interest.

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Authors’ contributions

AN and MSo participated in the design of the study, performed the statistical analysis and drafted the manuscript. HT and YaA participated in the design of the study and performed statistical analyses. YoA, MSa and MY participated in the design of the study and helped to draft the manuscript. KY, JS, MN and KK collected patients’ clinical information and made a part of figures and tables. TS and KN conceived of the study, participated in its design and coordination and helped to revise the manuscript. All authors read and approved the final manuscript.

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Tables
Due to technical limitations, table 1 to 6 is only available as a download in the Supplemental Files section.