Introduction

Any problems involving the patella after total knee arthroplasty (TKA) can influence overall knee function\(^1\). In particular, periprosthetic patellar fractures, albeit infrequent, have been considered a troublesome complication because return to pre-fracture function is rare even with anatomic reduction, healing, and reconstitution of the extensor mechanism\(^1-3\). It is known that the thickness of the residual patella is one of the most important factors affecting the risk of periprosthetic patellar fractures. Many previous studies have proposed that if the resurfaced patella is too thin, there may be an increased risk of fracture due to increased stress on the residual native bone\(^4-6\).

Many manufacturers have modified their TKA prosthesis designs to improve functional outcomes. The Attune prosthesis (Depuy Synthes, Warsaw, IN, USA) is a modified version of the PFC Sigma (Depuy Synthes). The theoretical advantages of the Attune prosthesis are increased conformity between the femoral component and polyethylene insert, an extensive range of sizes for diverse populations, optimization of patellofemoral conformity, and an improved polyethylene insert locking mechanism (Fig. 1)\(^7-10\). These design modifications resulted in improvement in patellofemoral clinical outcomes, decreasing the incidence of anterior knee pain and patellar crepitus\(^8-10\). However, despite such advantages, there are some design features of the current prosthesis that might cause problems. The thickness of the Attune patellar component is greater than that of the PFC Sigma. The thicknesses of the Attune patellar components are 8.5, 9.0, 9.5, 10, and 10.5 mm for sizes 29, 32, 35, 38, and 41, respectively\(^7\). By contrast, the thicknesses of the PFC Sigma...

Comparison of Clinical Results and Risk of Patellar Injury between Attune and PFC Sigma Knee Systems

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Purpose: The purposes of this study were to compare clinical results after total knee arthroplasty (TKA) using the Attune and PFC Sigma knee designs and to investigate whether the use of the Attune prosthesis increased the risk of patellar injury in Asian patients.

Materials and Methods: Three hundred knees that underwent TKA using Attune (group A) were compared to 300 knees that underwent TKA using PFC Sigma (group B). The Knee Society Knee Score (KS) and Function Score (FS), and range of motion (ROM) were compared. The residual patellar thickness was compared to evaluate the risk of patellar injury.

Results: The postoperative KS and ROM of group A were better than those of group B (93.1 vs. 88.8, \(p<0.001\) and 131.4° vs. 129.0°, \(p=0.008\), respectively). The postoperative FS did not differ significantly between the two groups (80.9 vs. 78.7, \(p=0.427\)). The residual patella was thinner in group A (14.8 mm vs. 15.7 mm, \(p=0.003\)), which made up a higher proportion of the high-risk group for patellar fractures with a residual thickness of <12 mm (7.5% vs. 2.1%, \(p=0.003\)).

Conclusions: TKA using the Attune prosthesis provided more favorable clinical results than TKA using PFC Sigma. However, the risk of injury in the residual patella was increased with use of the Attune prosthesis in Asian patients.

Keywords: Knee, Arthroplasty, Patella, Injury, Risk
patellar components are 8.0 mm for size 29; 8.5 mm for sizes 32, 35, and 38; and 11 mm for size 41\(^{(11)}\). Thus, considering that the postoperative patellar-patellar component composite thickness should be similar with the thickness of the original patella, the residual bone stock should be shallow in the knees with the current prosthesis, which increases the possibility of patellar fractures especially in patients with a small patella\(^{(12,13)}\). In general, Asians have a smaller knee with a thinner patella compared to Caucasian; therefore, the risk of patellar injury might be increased when the Attune prosthesis is used in Asian patients\(^{(12)}\).

The purposes of this study were to compare clinical and radiographic results after TKA using the Attune and PFC Sigma knee systems and to investigate whether the use of Attune prosthesis increased injury risk to the patella in Asian patients. It was hypothesized that the clinical results of Attune prosthesis would be comparable to or better than those of the PFC Sigma prosthesis and that the injury risk to the patella could be increased with use of Attune.

### Materials and Methods

#### 1. Patients

All consecutive patients who underwent TKA using Attune between November 2014 and June 2015 (group A) were enrolled in this study. During this period, 300 TKAs were performed with this prosthesis in 273 patients. For each patient reviewed, we matched a control from our patient database who had undergone primary TKA with PFC Sigma (group B) with respect to age, gender, body mass index, diagnosis, preoperative range of motion (ROM), and severity of preoperative deformity. The inclusion criterion was TKA performed by two senior surgeons with more than 20 years of experience in primary posterior stabilized TKA using the two prostheses. All patients underwent patella resurfacing due to severe patellofemoral arthritis. We excluded (1) patients with previous infection, fracture, or dislocation of the knee; (2) patients with previous high tibial osteotomy or revision TKA; and (3) patients with extra-articular deformity or severe bone loss. These patients were excluded because of the concern for anatomical distortion of the patella. The preoperative demographic data did not differ significantly between the two groups except the follow-up period (Table 1). The follow-up period was inevitably longer in group B because we have mainly used the Attune prosthesis instead of its predecessor (PFC Sigma) after introduction of the latest model. This study was approved by the Institutional Review Board (KHUH 2017-05-062). Informed consent was obtained from all patients before commencing the database review.

#### 2. Surgical Technique and Rehabilitation

The medial parapatellar approach was used with a midline skin incision. All TKAs were planned to use a posterior stabilized prosthesis with patella resurfacing. Femoral and tibial bone resections were made with a modified measured resection technique. The transepicondylar axis was used as a reference for femoral component rotation. The tibial resection was set to be 0°–3° of the posterior slope in the sagittal plane. The reference line for tibial rotation was accurately aimed to pass through the medial third of the tibial tubercle and the second metatarsal bone or the middle of the talus. All osteophytes were removed. Any contracted medial or lateral soft tissue was carefully evaluated and selectively released where required.

To prepare the patella, the patella was resected with a sharp
electric saw. For the evaluation of the even cut surface, the patella was divided into four quadrants and thickness was measured at the center of each quadrant using a caliper. The original patellar thickness was preserved or slightly decreased by about 0.5 mm with beveling of the uncovered lateral cut surface. The medialized dome patellar component was used for group A, and the oval dome patellar component was used for group B. Patellofemoral articulation was carefully evaluated with the no thumb technique. Lateral retinacular release was not performed because there was no case with patellar maltracking under tourniquet inflation.

The postoperative rehabilitation protocol was similar between the two groups\textsuperscript{14}. Isometric exercises using the extensor and flexor muscles were initiated shortly after operation. Drain was removed on the second postoperative day, followed by the initiation of active and assisted ROM exercises. Full weight bearing ambulation was started 4 days postoperatively to the extent that the patient’s condition permitted.

### 3. Clinical and Radiographic Evaluation

The clinical scores recorded in our database were retrospectively analyzed. For the clinical evaluation, Knee Society Knee Score (KS) and Function Score (FS) were used to access pain and function\textsuperscript{15}. The ROM was measured using a long-armed goniometer. Follow-up was conducted regularly at postoperative 3 months, 6 months, 1 year, and annually thereafter.

Pre- and postoperative anteroposterior (AP) and lateral radiographs and orthoreonentrionograms (full-length standing AP radiographs) were used to assess limb alignment and component positioning. Pre- and postoperative mechanical axis was defined as the angle between the femoral and tibial mechanical axes on orthorothionentograms. Detailed analyses of the AP and lateral radiographs were conducted to evaluate the position of components with $\alpha$, $\beta$, $\gamma$, and $\delta$ angles using the Knee Society radiological evaluation method (Fig. 2)\textsuperscript{16}.

The pre- and postoperative patella thickness and postoperative thickness of the residual patella were measured in the Merchant view of the knee\textsuperscript{17}. Patellar thickness was defined as the length of the thickest portion of the preoperative patella or postoperative prosthesis-patellar composite. The thickness of the residual patella was measured at the thickest portion of the remaining pa-

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### Table 1. Comparison of Patient Demographics between Groups

| Variable                  | Group A\textsuperscript{a} | Group B\textsuperscript{b} | p-value |
|---------------------------|-----------------------------|-----------------------------|---------|
| Operating period          | Nov 2014–Jun 2015           | Jan 2013–Nov 2014           |         |
| No. of knees (patients)   | 300 (273)                   | 300 (282)                   |         |
| Age (yr)                  | 69.7±7.7                    | 68.9±6.9                    | 0.102   |
| Sex (female/male)         | 287/13                      | 290/10                      | 0.524   |
| Right/left                | 151/149                     | 154/146                     | 0.806   |
| Body mass index (kg/m\textsuperscript{2}) | 26.5±3.6                   | 26.1±3.3                    | 0.565   |
| OA/RA/Others              | 289/5/6                     | 290/6/4                     | 0.455   |
| Range of motion (°)       | 119.8±25.7                  | 120.2±20.7                  | 0.834   |
| Preoperative mechanical axis (°) | Varus 11.6±6.6               | Varus 11.7±7.5               | 0.751   |
| Follow-up period (mo)     | 24.8±6.0 (16–37)            | 33.3±9.0 (21–76)            | <0.001  |

Values are presented as mean±standard deviation (range).

OA: osteoarthritis, RA: rheumatoid arthritis, Others: post-traumatic arthritis, infection sequelae, and hemophilic arthritis.

\textsuperscript{a}Group A patients who received the Attune prosthesis.

\textsuperscript{b}Group B patients who received the PFC Sigma prosthesis.
tella after resurfacing (Fig. 3). Because a patella fracture is more likely to develop when the residual thickness is less than 12 mm, patients with such thicknesses were categorized into a high-risk group for patellar fractures\textsuperscript{18,19}.

The quality of radiographic evaluation could be improved by standardization of the radiographic protocol with respect to the position of the knee and the distance between the X-ray beam and the cassette. The AP radiographs and orthoroentgenograms were taken with the patient standing with the knee fully extended and the feet slightly internally rotated to ensure forward placement of all anatomic landmarks\textsuperscript{20}. For the lateral radiographs, the knee was positioned in the same manner except the x-ray beam was directed laterally at 90° to the AP plane\textsuperscript{21}. The distance between the radiographic source and the patient’s bone was 245 cm, and that between the radiographic source and the cassette film was 260 cm\textsuperscript{22}.

When taking the Merchant radiographs, a standardized protocol was also used to minimize variances in the quality of radiographs. A leg position device equipped with an X-ray cassette holding slot was used to maintain uniformity in the angle and distances between the X-ray tube, the patella, and the cassette. The patient lay supine on the table with the knee flexed and supported at 45° angle. The x-ray beam was directed toward the feet at 30° from the horizontal, and a film cassette set was positioned 30 cm below the knee. The angle and distance between the X-ray source, the patella, and the cassette were kept unchanged.

The images were transferred digitally to a picture archiving and communication system (PACS; Infinitt, Seoul, Korea). Assessment was performed on a 61-cm (24-inch) monitor (SyncMaster 2494HMN; Samsung, Seoul, Korea) in portrait mode using the PACS software. The minimum differences that the software could detect were 0.1° in angle and 0.1 mm in length\textsuperscript{23}.

The radiographic evaluation of the Merchant view was performed on the PACS image setting the PACS ruler to be 10 cm. Magnification ratio was evaluated through a pilot test using 30 knees, and it was found to be approximately 1.1 in the PACS image. The measurements on the Merchant view were adjusted taking into account the magnification ratio.

To minimize any observation bias, two independent investigators repeatedly performed all of the radiographic measurements with an interval of 2 weeks. The intra- and interobserver reliabilities of all measurements were assessed using the intraclass correlation coefficient, all values of which were greater than 0.8. Thus, the radiographic measurements taken by one investigator who had more clinical experience than the other investigator were used in the analyses.

The actual occurrences of complications associated with the injury of patella, including periprosthetic fracture and loosening of the patellar component, were investigated during the follow-up period.

4. Statistical Analysis

Clinical and radiographic results were compared between groups A and B (independent Student \textit{t}-test). The preoperative and postoperative clinical and radiographic results were also compared (paired \textit{t}-test). The proportion of patients in the high-risk group for patella fracture was compared between the two groups (chi-square test).

To determine whether our sample had sufficient power to detect significant differences, we performed post-hoc power analyses using the significance levels set at an alpha of 0.05. A power >80% was considered sufficient, and all of the variables that were significantly different met the criterion. Thus, we determined that our study was adequately powered.

Statistical analyses were performed using SPSS ver. 18.0 (SPSS Inc., Chicago, IL, USA), and a \textit{p}<0.05 was considered statistically significant.

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Fig. 3. Radiographic measurement of the thickness of the original and residual patella. (A) Original patella. (B) Attune. (C) PFC Sigma. O: original patellar thickness, R: residual patellar thickness.
Results

The KS and FS improved significantly in both groups after TKA (p<0.001) (Table 2). The postoperative KS was higher in group A (93.1 vs. 88.8, p<0.001). The postoperative ROM of group A was also greater than that of group B (131.4° vs. 129.0°, p=0.008).

Radiographically, there were no significant differences in the pre- and postoperative mechanical axis between group A and group B (Table 3). The mean α, β, γ, and δ angles did not differ significantly between the two groups (Table 3).

There were no significant differences in pre- and postoperative patellar thickness between two groups (Table 3). The changes in patella thickness also did not differ significantly between group A and group B (0.7 mm vs. 0.6 mm, p=0.799). The residual patella was thinner in group A than group B (14.8 mm vs. 15.7 mm, p<0.001) (Table 3). Group A had more patients at high-risk for patellar fractures (7.5% vs. 2.1%, p=0.003) (Table 4).

Regarding the actual occurrence of complications, there was no knee in which a periprosthetic fracture or loosening around the patellar component occurred.

Discussion

The most important finding of the present study is that the risk of injury to residual patella after TKA might be higher when using Attune, although the prosthesis provided more favorable clinical results than PFC Sigma.

Several previous studies compared the clinical results between

| Table 2. Comparison of the Clinical Results between the Groups |
|------------------------|---------------------|---------------------|-----------------|
| Variable               | Group A             | Group B             | p-value         |
| Knee score             |                     |                     |                 |
| Preoperative           | 42.2±16.9           | 43.9±9.0            | 0.145           |
| Postoperative          | 93.1±6.4            | 88.8±6.2            | <0.001          |
| Change                 | 50.9±18.4           | 44.9±10.6           | <0.001          |
| Function score         |                     |                     |                 |
| Preoperative           | 43.6±13.5           | 40.2±13.6           | 0.002           |
| Postoperative          | 80.9±12.4           | 78.7±10.9           | 0.427           |
| Change                 | 37.3±15.4           | 38.5±15.3           | 0.201           |
| Range of motion (°)    |                     |                     |                 |
| Preoperative           | 119.8±25.7          | 120.2±20.7          | 0.834           |
| Postoperative          | 131.4±10.1          | 129.0±12.2          | 0.008           |
| Change                 | 11.6±24.2           | 8.8±19.5            | 0.113           |

Values are presented as mean±standard deviation.

a) Group A patients who received the Attune prosthesis.
b) Group B patients who received the PFC Sigma prosthesis.

| Table 3. Comparison of the Radiographic Results between the Groups |
|------------------------|---------------------|---------------------|-----------------|
| Variable               | Group A             | Group B             | p-value         |
| Mechanical axis (°)    |                     |                     |                 |
| Preoperative           | –11.6±6.6           | –11.7±7.5           | 0.751           |
| Postoperative          | –0.9±2.6            | –1.2±2.8            | 0.101           |
| Change                 | 10.7±6.6            | 10.5±7.1            | 0.145           |
| Position of components (°) |               |                     |                 |
| α angle                | 95.4±1.7            | 95.0±1.6            | 0.102           |
| β angle                | 90.8±2.1            | 90.6±2.2            | 0.158           |
| γ angle                | 2.3±2.9             | 1.5±2.3             | 0.091           |
| δ angle                | 88.7±2.5            | 88.5±2.1            | 0.101           |
| Patellar thickness (mm) |                     |                     |                 |
| Preoperative           | 23.1±3.0            | 23.6±3.0            | 0.088           |
| Postoperative          | 23.9±2.3            | 24.2±2.5            | 0.073           |
| Change                 | 0.7±2.9             | 0.6±3.4             | 0.799           |
| Thickness of the residual patella (mm) |     |                     |                 |
| Preoperative           | 14.8±2.1            | 15.7±2.4            | <0.001          |

Values are presented as mean±standard deviation.

a) Group A patients who received the Attune prosthesis.
b) Group B patients who received the PFC Sigma prosthesis.

c) The proportion of high-risk cases with a residual patellar thickness of less than 12 mm was significantly different between the groups (p=0.003).

Table 4. Comparison of the Distribution of Residual Patellar Thickness

| Thickness (mm) | Group A | Group B |
|---------------|---------|---------|
| 12–14         | 88      | 66      |
| 14–16         | 115     | 122     |
| 16–18         | 52      | 57      |
| 18–20         | 16      | 36      |
| 20–22         | 6       | 9       |
| 24–26         | 0       | 2       |
| 26–28         | 0       | 0       |
| 28–30         | 0       | 2       |

a) Thickness of the residual patella.
b) Group A patients who received the Attune prosthesis.
c) Group B patients who received the PFC Sigma prosthesis.
d) The proportion of high-risk cases with a residual patellar thickness of less than 12 mm was significantly different between the groups (p=0.003).

Attune and PFC sigma TKAs. Ranawat et al. reported 2-year follow-up clinical results based on Knee Society scores in groups with the above two prostheses: there were no significant differences between the two groups in the postoperative KS (92.4 vs. 92.8, p=0.75) and FS (89.3 vs. 89.4, p=0.096), and no difference
was also shown in the postoperative ROM (117.0° vs. 114.2°, p=0.025). However, in our study, there were better clinical results in the group with Attune with respect to the postoperative KS and ROM. These better results in group A might be associated with improvement in the design that allows for gradual reduction of the femoral radius during knee flexion from 5° to 70° and optimization of patellofemoral conformity.

Many previous studies reported that the risk of periprosthetic patellar fracture might increase when the residual patella is too thin. Reuben et al. evaluated the strain on the patellofemoral joint in a cadaveric study and concluded that patellar strain was increased when the thickness of the residual patella was less than 15 mm. A clinical study by Seo et al. evaluated the various risk factors that might be associated with patellar fractures after TKA. In the review of consecutive 7,866 TKAs, they found that patients with a patellar thickness of less than 12 mm had a greater risk of patellar fractures (odds ratio, 1.6; p<0.043). Although, some literatures had conflicting views on the role of the residual patella in small knees with a preoperative patellar thickness of <20 mm. Asian patients, especially when the Attune prosthesis is implanted, a greater precaution will be necessary to avoid patellar fractures in small knees with a preoperative patellar thickness of <12 mm (7.5% vs. 2.1%, p=0.003).

The risk of patellar fractures can be linked to the bony geometric characteristics of Asian people. The patella tends to be thinner in Asians. Kim et al. reported that the average thickness of the patella in Koreans is 21.2 mm in females and 23.1 mm in males, whereas the corresponding mean thickness in Caucasians is 21.8 to 22.5 mm in females and 23.9 to 26.1 mm in males. Therefore, greater precaution will be necessary to avoid patellar fractures in Asian patients, especially when the Attune prosthesis is implanted in small knees with a preoperative patellar thickness of <20 mm.

After recognizing the risk of patellar injury in the Attune prosthesis, the residual patellar thickness has been advised to be at least 12 mm when using the latest prosthesis, even in the situation where the postoperative patella is expected to be thicker than the preoperative patella. The patella has been rarely overstuffed because the anterior flange of the current prosthesis is shallower than that of the previous one. If patellar overstuffing is expected, it could be prevented by displacing the femoral component posteriorly to the extent that notching of the anterior femoral cortex or excessive narrowing of the flexion gap does not occur.

This study has several limitations attributable to the retrospective design and relatively short follow-up duration. First, we assessed the injury risk based on radiographic measurements instead of the evaluation of actual complications after a long-term follow-up. The average 2-year follow-up could be sufficient for evaluation of radiographic parameters for the patellar thickness, not for the investigation of the occurrence of postoperative complications. Furthermore, regarding the low incidence rate of patellar fractures after TKA (about 1%), the number of cases in each group does not seem to be sufficient for proper intergroup comparison. Therefore, a larger cohort study with a longer follow up will be required. Second, cartilage thickness was not taken into consideration in the evaluation of the patellar thickness. This is the reason why the change of patellar thickness had a positive value. However, we followed the surgical principle that the original patellar thickness should be preserved or slightly decreased by 0.5 mm. In addition, measurement of the cartilage thickness was not considered necessary because the critical parameter of the present study was the residual patellar thickness. Third, most of the patients in the present study were Korean females with small knees, which should be considered when extrapolating our findings to other populations. However, based on our findings, we would like to emphasize the need for caution when using the Attune prosthesis in Asian patients to avoid the risk of injury to the residual patella.

Conclusions

The Attune knee system provided slightly better clinical results than the PFC Sigma prosthesis in TKA. However, the injury risk to the residual patella was increased with use of the Attune prosthesis in Asian patients.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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